



REPUBLIC OF KENYA

THIRTEENTH PARLIAMENT – (THIRD SESSION)

THE NATIONAL ASSEMBLY

ORDERS OF THE DAY

SUPPLEMENTARY

THURSDAY, JUNE 6, 2024 AT 2.30 P.M.

ORDER OF BUSINESS

PRAYERS

1. Administration of Oath
2. Communication from the Chair
3. Messages
4. Petitions
5. Papers
6. Notices of Motion
7. Questions and Statements

8*. PROCEDURAL MOTION – EXTENSION OF SITTING UNTIL CONCLUSION OF SPECIFIED BUSINESS

(The Leader of the Majority Party)

THAT, pursuant to the provisions of Standing Order 30(3)(a), this House **orders that**, should the time appointed for adjournment of the House be reached before conclusion of business under Order No. 10 (*Committee of Supply*), the sitting shall stand extended until conclusion of the particular business.

9*. MOTION – REPORT OF THE COMMITTEE OF THE WHOLE HOUSE ON THE KENYA DRUGS AUTHORITY BILL (NATIONAL ASSEMBLY BILL NO. 54 OF 2022)

(The Hon. (Dr.) Robert Pukose, M.P.)

THAT, this House do agree with the Report of the Committee of the Whole House on its consideration of the Kenya Drugs Authority Bill (National Assembly Bill No. 54 of 2022) up to clause 54 and seeks leave to sit again.

(*Question to be put*)

10*. COMMITTEE OF SUPPLY

Estimates of Recurrent and Development Expenditure for the year ending 30th June 2025

IN THE COMMITTEE
THE EXECUTIVE

Vote 1011 - Executive Office of the President

THAT, a sum not exceeding **Kshs. 5,132,190,119** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1011 (Executive Office of the President)**.

Vote 1012 - Office of the Deputy President

THAT, a sum not exceeding **Kshs. 4,896,000,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1012 (Office of the Deputy President)**.

Vote 1013 - Office of the Prime Cabinet Secretary

THAT, a sum not exceeding **Kshs. 1,140,788,324** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1013 (Office of the Prime Cabinet Secretary)**.

Vote 1014 - State Department for Parliamentary Affairs

THAT, a sum not exceeding **Kshs. 458,283,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1014 (State Department for Parliamentary Affairs)**.

Vote 1015 - State Department for Performance and Delivery Management

THAT, a sum not exceeding **Kshs. 597,112,861** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1015 (State Department for Performance and Delivery Management)**.

Vote 1016 - State Department for Cabinet Affairs

THAT, a sum not exceeding **Kshs. 275,136,014** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1016 (State Department for Cabinet Affairs)**.

Vote 1017 - State House

THAT, a sum not exceeding **Kshs. 9,496,000,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1017 (State House)**.

Vote 1023 - State Department for Correctional Services

THAT, a sum not exceeding **Kshs. 35,550,346,616** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1023 (State Department for Correctional Services)**.

Vote 1024 - State Department for Immigration and Citizen Services

THAT, a sum not exceeding **Kshs. 14,959,813,872** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1024 (State Department for Immigration and Citizen Services)**.

Vote 1025 - National Police Service

THAT, a sum not exceeding **Kshs. 110,552,072,775** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1025 (National Police Service)**.

Vote 1026 - State Department for Internal Security and National Administration

THAT, a sum not exceeding **Kshs. 35,877,114,720** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1026 (State Department for Internal Security and National Administration)**.

Vote 1032 - State Department for Devolution

THAT, a sum not exceeding **Kshs. 4,242,428,367** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1032 (State Department for Devolution)**.

Vote 1036 - State Department for ASALs and Regional Development

THAT, a sum not exceeding **Kshs. 12,540,387,586** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1036 (State Department for ASALs and Regional Development)**.

Vote 1041 - Ministry of Defence

THAT, a sum not exceeding **Kshs. 173,086,817,170** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1041 (Ministry of Defence)**.

Vote 1053 - State Department for Foreign Affairs

THAT, a sum not exceeding **Kshs. 23,097,447,602** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1053 (State Department for Foreign Affairs)**.

Vote 1054 - State Department for Diaspora Affairs

THAT, a sum not exceeding **Kshs. 828,143,693** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1054 (State Department for Diaspora Affairs)**.

Vote 1064 - State Department for Technical, Vocational Education and Training

THAT, a sum not exceeding **Kshs. 30,688,638,919** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1064 (State Department for Technical, Vocational Education and Training)**.

Vote 1065 - State Department for Higher Education and Research

THAT, a sum not exceeding **Kshs. 127,986,194,444** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1065 (State Department for Higher Education and Research)**.

Vote 1066 - State Department for Basic Education

THAT, a sum not exceeding **Kshs. 142,263,122,192** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1066 (State Department for Basic Education)**.

Vote 1071 - The National Treasury

THAT, a sum not exceeding **Kshs. 146,072,137,600** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1071 (The National Treasury)**.

Vote 1072 - State Department for Economic Planning

THAT, a sum not exceeding **Kshs. 66,782,133,355** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1072 (State Department for Economic Planning)**.

Vote 1082 - State Department for Medical Services

THAT, a sum not exceeding **Kshs. 102,983,407,719** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1082 (State Department for Medical Services)**.

Vote 1083 - State Department for Public Health and Professional Standards

THAT, a sum not exceeding **Kshs. 28,037,735,123** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1083 (State Department for Public Health and Professional Standards)**.

Vote 1091 - State Department for Roads

THAT, a sum not exceeding **Kshs. 199,361,988,075** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1091 (State Department for Roads)**.

Vote 1092 - State Department for Transport

THAT, a sum not exceeding **Kshs. 51,631,203,728** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1092 (State Department for Transport)**.

Vote 1093 - State Department for Shipping and Maritime Affairs

THAT, a sum not exceeding **Kshs. 3,593,974,935** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1093 (State Department for Shipping and Maritime Affairs)**.

Vote 1094 - State Department for Housing and Urban Development

THAT, a sum not exceeding **Kshs. 87,566,192,681** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1094 (State Department for Housing and Urban Development)**.

Vote 1095 - State Department for Public Works

THAT, a sum not exceeding **Kshs. 5,009,078,552** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1095 (State Department for Public Works)**.

Vote 1104 - State Department for Irrigation

THAT, a sum not exceeding **Kshs. 23,382,162,500** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1104 (State Department for Irrigation)**.

Vote 1109 - State Department for Water and Sanitation

THAT, a sum not exceeding **Kshs. 52,409,738,911** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1109 (State Department for Water and Sanitation)**.

Vote 1112 - State Department for Lands and Physical Planning

THAT, a sum not exceeding **Kshs. 8,770,536,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1112 (State Department for Lands and Physical Planning)**.

**Vote 1122 - State Department for Information Communication
Technology and Digital Economy**

THAT, a sum not exceeding **Kshs. 22,354,880,752** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1122 (State Department for Information Communication Technology and Digital Economy)**.

**Vote 1123 - State Department for Broadcasting and
Telecommunications**

THAT, a sum not exceeding **Kshs. 6,061,310,364** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1123 (State Department for Broadcasting and Telecommunications)**.

Vote 1132 - State Department for Sports

THAT, a sum not exceeding **Kshs. 17,743,686,404** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1132 (State Department for Sports)**.

Vote 1134 - State Department for Culture and Heritage

THAT, a sum not exceeding **Kshs. 3,030,997,321** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1134 (State Department for Culture and Heritage)**.

Vote 1135 - State Department for Youth Affairs and Creative Economy

THAT, a sum not exceeding **Kshs. 4,048,471,229** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1135 (State Department for Youth Affairs and Creative Economy)**.

Vote 1152 - State Department for Energy

THAT, a sum not exceeding **Kshs. 69,663,515,207** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1152 (State Department for Energy)**.

Vote 1162 - State Department for Livestock Development

THAT, a sum not exceeding **Kshs. 12,316,954,089** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1162 (State Department for Livestock)**.

Vote 1166 - State Department for the Blue Economy and Fisheries

THAT, a sum not exceeding **Kshs. 11,996,725,869** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1166 (State Department for the Blue Economy and Fisheries)**.

Vote 1169 - State Department for Crop Development

THAT, a sum not exceeding **Kshs. 48,230,425,195** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1169 (State Department for Crop Development)**.

Vote 1173 - State Department for Cooperatives

THAT, a sum not exceeding **Kshs. 4,080,953,583** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1173 (State Department for Cooperatives)**.

Vote 1174 - State Department for Trade

THAT, a sum not exceeding **Kshs. 3,598,361,146** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1174 (State Department for Trade)**.

Vote 1175 - State Department for Industry

THAT, a sum not exceeding **Kshs. 8,439,676,621** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1175 (State Department for Industry)**.

Vote 1176 - State Department for Micro, Small and Medium Enterprises Development

THAT, a sum not exceeding **Kshs. 9,369,058,500** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1176 (State Department for Micro, Small and Medium Enterprises Development)**.

Vote 1177 - State Department for Investment Promotion

THAT, a sum not exceeding **Kshs. 4,616,043,914** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1177 (State Department for Investment Promotion)**.

Vote 1184 - State Department for Labour and Skills Development

THAT, a sum not exceeding **Kshs. 5,832,415,243** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1184 (State Department for Labour and Skills Development)**.

Vote 1185 - State Department for Social Protection and Senior Citizen Affairs

THAT, a sum not exceeding **Kshs. 35,304,805,645** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1185 (State Department for Social Protection and Senior Citizen Affairs)**.

Vote 1192 - State Department for Mining

THAT, a sum not exceeding **Kshs. 1,758,158,447** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1192 (State Department for Mining)**.

Vote 1193 - State Department for Petroleum

THAT, a sum not exceeding **Kshs. 31,200,411,883** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1193 (State Department for Petroleum)**.

Vote 1202 - State Department for Tourism

THAT, a sum not exceeding **Kshs. 10,328,821,808** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1202 (State Department for Tourism)**.

Vote 1203 - State Department for Wildlife

THAT, a sum not exceeding **Kshs. 14,343,194,935** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1203 (State Department for Wildlife)**.

Vote 1212 - State Department for Gender and Affirmative Action

THAT, a sum not exceeding **Kshs. 6,114,541,404** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1212 (State Department for Gender and Affirmative Action)**.

Vote 1213 - State Department for Public Service

THAT, a sum not exceeding **Kshs. 19,612,689,909** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1213 (State Department for Public Service)**.

Vote 1221 - State Department for East African Community

THAT, a sum not exceeding **Kshs. 647,487,899** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1221 (State Department for East African Community)**.

Vote 1252 - State Law Office

THAT, a sum not exceeding **Kshs. 6,977,570,997** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1252 (State Law Office)**.

Vote 1271 - Ethics and Anti-Corruption Commission

THAT, a sum not exceeding **Kshs. 4,171,550,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1271 (Ethics and Anti-Corruption Commission)**.

Vote 1281 - National Intelligence Service

THAT, a sum not exceeding **Kshs. 46,351,000,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1281 (National Intelligence Service)**.

Vote 1291 - Office of the Director of Public Prosecutions

THAT, a sum not exceeding **Kshs. 4,007,520,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1291 (Office of the Director of Public Prosecutions)**.

Vote 1311 - Office of the Registrar of Political Parties

THAT, a sum not exceeding **Kshs. 2,037,871,453** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1311 (Office of the Registrar of Political Parties)**.

Vote 1321 - Witness Protection Agency

THAT, a sum not exceeding **Kshs. 741,192,500** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1321 (Witness Protection Agency)**.

Vote 1331 - State Department for Environment and Climate Change

THAT, a sum not exceeding **Kshs. 5,019,131,295** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1331 (State Department for Environment and Climate Change)**.

Vote 1332 - State Department for Forestry

THAT, a sum not exceeding **Kshs. 12,563,930,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1332 (State Department for Forestry)**.

Vote 2011 - Kenya National Commission on Human Rights

THAT, a sum not exceeding **Kshs. 478,074,025** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2011 (Kenya National Commission on Human Rights)**.

Vote 2021 - National Land Commission

THAT, a sum not exceeding **Kshs. 2,016,222,679** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2021 (National Land Commission)**.

Vote 2031 - Independent Electoral and Boundaries Commission

THAT, a sum not exceeding **Kshs. 3,755,219,680** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2031 (Independent Electoral and Boundaries Commission)**.

Vote 2061 - The Commission on Revenue Allocation

THAT, a sum not exceeding **Kshs. 413,465,304** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2061 (The Commission on Revenue Allocation)**.

Vote 2071 - Public Service Commission

THAT, a sum not exceeding **Kshs. 3,667,530,017** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2071 (Public Service Commission)**.

Vote 2081 - Salaries and Remuneration Commission

THAT, a sum not exceeding **Kshs. 472,230,922** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2081 (Salaries and Remuneration Commission)**.

Vote 2091 - Teachers Service Commission

THAT, a sum not exceeding **Kshs. 358,216,066,118** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2091 (Teachers Service Commission)**.

Vote 2101 - National Police Service Commission

THAT, a sum not exceeding **Kshs. 1,131,272,317** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2101 (National Police Service Commission)**.

Vote 2111 - Auditor General

THAT, a sum not exceeding **Kshs. 8,666,770,850** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2111 (Auditor General)**.

Vote 2121 - Controller of Budget

THAT, a sum not exceeding **Kshs. 740,219,080** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2121 (Controller of Budget)**.

Vote 2131 - Commission on Administrative Justice

THAT, a sum not exceeding **Kshs. 661,974,500** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2131 (Commission on Administrative Justice)**.

Vote 2141 - National Gender and Equality Commission

THAT, a sum not exceeding **Kshs. 435,810,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2141 (National Gender and Equality Commission)**.

Vote 2151 - Independent Policing Oversight Authority

THAT, a sum not exceeding **Kshs. 1,107,672,060** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2151 (Independent Policing Oversight Authority)**.

THE JUDICIARY

Vote 1261 - The Judiciary

THAT, a sum not exceeding **Kshs. 23,737,400,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 1261 (The Judiciary)**.

Vote 2051 - Judicial Service Commission

THAT, a sum not exceeding **Kshs. 902,900,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2051 (Judicial Service Commission)**.

PARLIAMENT

Vote 2041 - Parliamentary Service Commission

THAT, a sum not exceeding **Kshs. 1,167,000,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2041 (Parliamentary Service Commission)**.

Vote 2042 - National Assembly

THAT, a sum not exceeding **Kshs. 26,775,000,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2042 (National Assembly)**.

Vote 2043 - Parliamentary Joint Services

THAT, a sum not exceeding **Kshs. 8,646,000,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2043 (Parliamentary Joint Services)**.

Vote 2044 - Senate

THAT, a sum not exceeding **Kshs. 8,010,000,000** be issued from the Consolidated Fund to complete the sum necessary to meet the expenditure (Recurrent and Development) during the year ending 30th June, 2025 in respect of **Vote 2044 (Senate)**.

11*. COMMITTEE OF THE WHOLE HOUSE

The Kenya Drugs Authority Bill (National Assembly Bill No. 54 of 2022)
(The Hon. (Dr.) Robert Pukose, M.P)

(To resume from Clause 55)

12*. THE STATUTE LAW (MISCELLANEOUS AMENDMENTS) BILL
(NATIONAL ASSEMBLY BILL NO. 67 OF 2023)

(The Leader of the Majority Party)

Second Reading

(Resumption of debate interrupted on Thursday, June 06, 2024 - Morning Sitting)

13*. THE LAND LAWS (AMENDMENT) (No. 2) BILL (NATIONAL ASSEMBLY BILL NO. 76 OF 2023)

(The Leader of the Majority Party)

Second Reading

14*. MOTION – REPORTS OF THE AUDITOR-GENERAL ON THE NATIONAL GOVERNMENT CONSTITUENCIES DEVELOPMENT FUND FOR NINE CONSTITUENCIES IN BUNGOMA COUNTY

(The Chairperson, Decentralized Funds Accounts Committee)

THAT, this House **adopts** the Report of the Decentralized Funds Accounts Committee on its consideration of the Reports of the Auditor-General on the Financial Statements for the National Government Constituencies Development Fund for nine Constituencies in Bungoma County for Financial Years 2013/2014, 2014/2015 and 2015/2016, *laid on the Table of the House on Thursday, 7th March 2024.*

15*. MOTION – PUBLIC PETITION ON FUNDS SPENT CONTRARY TO THE PROVISIONS OF ARTICLE 223 OF THE CONSTITUTION

(The Chairperson, Public Petitions Committee)

THAT, this House **adopts** the Report of the Public Petitions Committee on its consideration of Public Petition No. 4 of 2022 regarding Funds Spent by the National Government Contrary to the Provisions of Article 223 of the Constitution, *laid on the Table of the House on Tuesday, 12th March 2024.*

16*. MOTION – REPORTS OF THE AUDITOR-GENERAL ON THE FINANCIAL STATEMENTS OF SPECIFIED STATE CORPORATIONS

(The Chairperson, Public Investments Committee on Social Services, Administration and Agriculture)

THAT, this House **adopts** the Second Report of the Public Investments Committee on Social Services, Administration and Agriculture on its consideration of the Reports of the Auditor-General on the Financial Statements of the Kenya Medical Training College for Financial Years 2017/2018 to 2020/2021; the Kenyatta University Teaching Referral and Research Hospital for Financial Years 2019/2020; the Kenya Medical Research Institute for Financial Years 2012/2013 and 2013/2014; the Kenya Medical Laboratory Technicians and Technologists Board for Financial Years 2014/2015 to 2020/2021; and the Kenya Medical Practitioners and Dentists Council for Financial Years 2017/2018 to 2020/2021, *laid on the Table of the House on Thursday, 22nd February 2024.*

17*. MOTION – ENHANCING REPORTING OF PARLIAMENTARY BUSINESS ON ONLINE PLATFORMS

(The Chairperson, Committee on Parliamentary Broadcasting and Library)

THAT, this House **adopts** the Report of the Select Committee on Parliamentary Broadcasting and Library on Enhancing Reporting of Parliamentary Business on Online Platforms, *laid on the Table of the House on Wednesday, 4th October 2023*.

18*. MOTION – INSPECTION VISIT TO SEMI-AUTONOMOUS INSTITUTIONS OF THE EAST AFRICAN COMMUNITY IN UGANDA

(The Chairperson, Committee on Regional Integration)

THAT, this House **adopts** the Report of the Committee on Regional Integration on its consideration of the Inspection Visit to the Semi-autonomous Institutions of the East African Community in Uganda, *laid on the Table of the House on Wednesday, 29th November 2023*.

19*. MOTION – IMPLEMENTATION STATUS OF REPORTS ON PETITIONS AND RESOLUTIONS PASSED BY THE HOUSE

(The Chairperson, Committee on Implementation)

THAT, this House **adopts** the Report of the Select Committee on Implementation on its consideration of the First Report on Implementation Status of Reports on Petitions and Resolutions passed by the House, *laid on the Table of the House on Thursday, 26th October 2023*.

20*. MOTION – ALLEGED UNFAIR TRADE PRACTICES BY FOREIGN INVESTORS IN KENYA

(The Chairperson, Departmental Committee on Trade, Industry and Cooperatives)

THAT, this House **adopts** the Report of the Departmental Committee on Trade, Industry and Cooperatives on the Inquiry into Alleged Unfair Trade Practices by Foreign Investors in Kenya, *laid on the Table of the House on Thursday, 7th March 2024*.

Denotes Orders of the Day

SCHEDULE**ANNUAL ESTIMATES FOR FY 2024/25 (IN KSHS)**

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25							
VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
1011	Executive Office of the President	4,521,144,412	852,300,000	5,373,444,412	4,231,290,119	900,900,000	5,132,190,119
	0603000 Government Printing Services	720,820,983	500,000,000	1,220,820,983	717,396,876	700,000,000	1,417,396,876
	0701000 General Administration Planning and Support Services	1,750,228,062	194,300,000	1,944,528,062	1,548,134,139	42,900,000	1,591,034,139
	0703000 Government Advisory Services	1,315,851,156	-	1,315,851,156	1,277,494,730	-	1,277,494,730
	Programme: Leadership and Coordination of Government Services	734,244,211	158,000,000	892,244,211	688,264,374	158,000,000	846,264,374
1012	Office of the Deputy President	4,575,600,000	300,400,000	4,876,000,000	4,575,600,000	320,400,000	4,896,000,000
	0734000 Deputy President Services	4,575,600,000	300,400,000	4,876,000,000	4,575,600,000	320,400,000	4,896,000,000
1013	Office of the Prime Cabinet Secretary	1,140,788,324	-	1,140,788,324	1,140,788,324	-	1,140,788,324
	0755000 Government Coordination and Supervision Services	1,140,788,324	-	1,140,788,324	1,140,788,324	-	1,140,788,324
1014	State Department for Parliamentary Affairs	458,283,000	-	458,283,000	458,283,000	-	458,283,000
	0759000 Parliamentary Liaison and Legislative Affairs	119,032,080	-	119,032,080	119,032,080	-	119,032,080

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	0760000 Policy Coordination and Strategy	117,042,480	-	117,042,480	117,042,480	-	117,042,480
	0761000 General Administration, Planning and Support Services	222,208,440	-	222,208,440	222,208,440	-	222,208,440
	State Department for Performance and Delivery Management	607,112,861	-	607,112,861	597,112,861	-	597,112,861
	0762000 Public Service Performance Management and Delivery Services	115,149,779	-	115,149,779	105,149,779	-	105,149,779
1015	0764000 General Administration, Planning and Support Services	218,111,458	-	218,111,458	218,111,458	-	218,111,458
	0772000 Service Delivery Management	228,523,344	-	228,523,344	228,523,344	-	228,523,344
	077300 Coordination and Supervision of Government Services	45,328,280	-	45,328,280	45,328,280	-	45,328,280
1016	State Department for Cabinet Affairs	275,136,014	-	275,136,014	275,136,014	-	275,136,014
	0758000 Cabinet Affairs Services	275,136,014	-	275,136,014	275,136,014	-	275,136,014
1017	State House	7,937,300,000	1,558,700,000	9,496,000,000	7,937,300,000	1,558,700,000	9,496,000,000
	0704000 State House Affairs	7,937,300,000	1,558,700,000	9,496,000,000	7,937,300,000	1,558,700,000	9,496,000,000
	State Department for Correctional Services	34,727,321,616	823,025,000	35,550,346,616	34,727,321,616	823,025,000	35,550,346,616
1023	0623000 General Administration, Planning and Support Services	603,685,214	7,000,000	610,685,214	603,685,214	7,000,000	610,685,214
	0627000 Prison Services	31,970,991,573	689,730,986	32,660,722,559	31,970,991,573	689,730,986	32,660,722,559

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	0628000 Probation & After Care Services	2,152,644,829	126,294,014	2,278,938,843	2,152,644,829	126,294,014	2,278,938,843
1024	State Department for Immigration and Citizen Services	10,289,613,872	4,861,000,000	15,150,613,872	10,149,613,872	4,810,200,000	14,959,813,872
	0605000 Migration & Citizen Services Management	4,404,014,189	2,635,000,000	7,039,014,189	4,334,014,189	2,635,000,000	6,969,014,189
	0626000 Population Management Services	4,751,107,352	2,136,000,000	6,887,107,352	4,751,107,352	2,085,200,000	6,836,307,352
	0631000 General Administration and Planning	1,134,492,331	90,000,000	1,224,492,331	1,064,492,331	90,000,000	1,154,492,331
1025	National Police Service	108,701,352,775	1,745,720,000	110,447,072,775	108,771,352,775	1,780,720,000	110,552,072,775
	0601000 Policing Services	108,701,352,775	1,745,720,000	110,447,072,775	108,771,352,775	1,780,720,000	110,552,072,775
1026	State Department for Internal Security & National Administration	27,831,624,720	7,155,290,000	34,986,914,720	28,311,624,720	7,565,490,000	35,877,114,720
	0629000 General Administration and Support Services	8,587,486,400	6,600,460,000	15,187,946,400	9,067,486,400	7,010,660,000	16,078,146,400
	0630000 Policy Coordination Services	1,343,357,100	65,000,000	1,408,357,100	1,343,357,100	65,000,000	1,408,357,100
	0632000 National Government Field Administration Services	17,900,781,220	489,830,000	18,390,611,220	17,900,781,220	489,830,000	18,390,611,220
1032	State Department for Devolution	1,489,428,367	2,653,000,000	4,142,428,367	1,589,428,367	2,653,000,000	4,242,428,367
	0712000 Devolution Services	1,489,428,367	2,653,000,000	4,142,428,367	1,589,428,367	2,653,000,000	4,242,428,367

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
1036	State Department for ASALs and Regional Development	4,169,293,586	5,907,894,000	10,077,187,586	4,857,493,586	7,682,894,000	12,540,387,586
	0733000 Accelerated ASAL Development	2,437,687,562	2,024,904,000	4,462,591,562	2,908,987,562	2,024,904,000	4,933,891,562
	0743000 General Administration, Planning and Support Services	378,489,218	-	378,489,218	478,489,218	-	478,489,218
	1013000 Integrated Regional Development	1,353,116,806	3,882,990,000	5,236,106,806	1,470,016,806	5,657,990,000	7,128,006,806
1041	Ministry of Defence	161,232,817,170	6,854,000,000	168,086,817,170	171,552,817,170	1,534,000,000	173,086,817,170
	0801000 Defence	157,961,800,000	6,854,000,000	164,815,800,000	168,281,800,000	1,534,000,000	169,815,800,000
	0802000 Civil Aid	350,000,000	-	350,000,000	350,000,000	-	350,000,000
	0803000 General Administration, Planning and Support Services	2,608,517,170	-	2,608,517,170	2,608,517,170	-	2,608,517,170
	0805000 National Space Management	312,500,000	-	312,500,000	312,500,000	-	312,500,000
1053	State Department for Foreign Affairs	20,707,347,602	2,390,100,000	23,097,447,602	20,707,347,602	2,390,100,000	23,097,447,602
	0714000 General Administration Planning and Support Services	2,767,407,779	318,000,000	3,085,407,779	2,767,407,779	318,000,000	3,085,407,779
	0715000 Foreign Relation and Diplomacy	17,746,053,022	1,966,890,000	19,712,943,022	17,746,053,022	1,966,890,000	19,712,943,022
	0741000 Economic and Commercial Diplomacy	49,859,065	-	49,859,065	49,859,065	-	49,859,065
	0742000 Foreign Policy Research, Capacity Development	144,027,736	105,210,000	249,237,736	144,027,736	105,210,000	249,237,736

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	and Technical Cooperation						
1054	State Department for Diaspora Affairs	828,143,693	-	828,143,693	828,143,693	-	828,143,693
	0752000 Management of Diaspora and Consular Affairs	828,143,693	-	828,143,693	828,143,693	-	828,143,693
1064	State Department for Technical Vocational Education and Training	23,039,038,919	6,927,000,000	29,966,038,919	23,129,038,919	7,559,600,000	30,688,638,919
	0505000 Technical Vocational Education and Training	22,544,742,427	6,927,000,000	29,471,742,427	22,634,742,427	7,559,600,000	30,194,342,427
	0507000 Youth Training and Development	58,918,193	-	58,918,193	58,918,193	-	58,918,193
	0508000 General Administration, Planning and Support Services	435,378,299	-	435,378,299	435,378,299	-	435,378,299
1065	State Department for Higher Education and Research	116,806,154,444	3,324,640,000	120,130,794,444	123,611,554,444	4,374,640,000	127,986,194,444
	0504000 University Education	115,711,858,039	3,110,640,000	118,822,498,039	122,677,258,039	4,310,640,000	126,987,898,039
	0506000 Research, Science, Technology and Innovation	836,497,147	214,000,000	1,050,497,147	636,497,147	64,000,000	700,497,147
	0508000 General Administration, Planning and Support Services	257,799,258	-	257,799,258	297,799,258	-	297,799,258
1066	State Department for Basic Education	119,937,562,192	19,203,560,000	139,141,122,192	121,927,562,192	20,335,560,000	142,263,122,192

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	0501000 Primary Education	12,893,724,959	12,785,000,000	25,678,724,959	16,393,724,959	13,542,000,000	29,935,724,959
	0502000 Secondary Education	98,419,234,354	6,337,960,000	104,757,194,354	96,619,234,354	6,722,960,000	103,342,194,354
	0503000 Quality Assurance and Standards	4,105,175,684	53,000,000	4,158,175,684	4,255,175,684	53,000,000	4,308,175,684
	0508000 General Administration, Planning and Support Services	4,519,427,195	27,600,000	4,547,027,195	4,659,427,195	17,600,000	4,677,027,195
1071	The National Treasury	66,992,982,220	61,993,955,380	128,986,937,600	83,245,982,220	62,826,155,380	146,072,137,600
	0717000 General Administration Planning and Support Services	57,157,494,314	7,709,499,380	64,866,993,694	73,621,194,314	10,941,699,380	84,562,893,694
	0718000 Public Financial Management	8,253,061,087	44,530,082,000	52,783,143,087	8,015,061,087	42,130,082,000	50,145,143,087
	0719000 Economic and Financial Policy Formulation and Management	1,073,922,429	9,754,374,000	10,828,296,429	1,101,222,429	9,754,374,000	10,855,596,429
	0720000 Market Competition	508,504,390	-	508,504,390	508,504,390	-	508,504,390
1072	State Department for Economic Planning	2,991,893,355	61,474,940,000	64,466,833,355	2,941,893,355	63,840,240,000	66,782,133,355
	0710000 Public Service Transformation	2,000,000	-	2,000,000	2,000,000	-	2,000,000
	07710000 Monitoring and Evaluation Services	112,200,000	7,740,000	119,940,000	112,200,000	7,740,000	119,940,000
	0706000 Economic Policy and National Planning	1,776,374,273	60,160,750,000	61,937,124,273	1,926,374,273	63,226,050,000	65,152,424,273
	0707000 National Statistical Information Services	867,848,000	1,306,450,000	2,174,298,000	867,848,000	606,450,000	1,474,298,000

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	0709000 General Administration Planning and Support Services	233,471,082	-	233,471,082	33,471,082	-	33,471,082
1082	State Department for Medical Services	60,496,874,385	39,516,533,334	100,013,407,719	68,096,874,385	34,886,533,334	102,983,407,719
	0402000 National Referral & Specialized Services	43,347,164,861	20,975,666,667	64,322,831,528	45,012,164,861	16,465,666,667	61,477,831,528
	0410000 Curative & Reproductive Maternal New Born Child Adolescent Health RMNCAH	1,379,140,420	15,354,866,667	16,734,007,087	1,479,140,420	15,434,866,667	16,914,007,087
	0411000 Health Research and Innovations	2,790,450,000	1,137,000,000	3,927,450,000	3,125,450,000	937,000,000	4,062,450,000
	0412000 General Administration	12,980,119,104	2,049,000,000	15,029,119,104	18,480,119,104	2,049,000,000	20,529,119,104
1083	State Department for Public Health and Professional Standards	21,781,755,123	5,053,180,000	26,834,935,123	22,473,555,123	5,564,180,000	28,037,735,123
	0406000 Preventive and Promotive Health Services	4,737,846,807	4,088,000,000	8,825,846,807	4,779,646,807	4,228,000,000	9,007,646,807
	0407000 Health Resources Development and Innovation	13,016,217,004	924,000,000	13,940,217,004	13,016,217,004	1,295,000,000	14,311,217,004
	0408000 Health Policy, Standards and Regulations	3,476,324,378	41,180,000	3,517,504,378	4,026,324,378	41,180,000	4,067,504,378
	0412000 General Administration	551,366,934	-	551,366,934	651,366,934	-	651,366,934
1091	State Department for Roads	72,086,891,250	112,416,896,825	184,503,788,075	72,176,891,250	127,185,096,825	199,361,988,075
	0202000 Road Transport	72,086,891,250	112,416,896,825	184,503,788,075	72,176,891,250	127,185,096,825	199,361,988,075

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
1092	State Department for Transport	15,613,803,728	44,935,400,000	60,549,203,728	16,397,803,728	35,233,400,000	51,631,203,728
	0201000 General Administration, Planning and Support Services	855,116,899	1,288,000,000	2,143,116,899	1,285,116,899	1,358,000,000	2,643,116,899
	0203000 Rail Transport	532,056,937	38,435,900,000	38,967,956,937	686,056,937	28,963,900,000	29,649,956,937
	0204000 Marine Transport	524,035,822	2,865,000,000	3,389,035,822	524,035,822	2,565,000,000	3,089,035,822
	0205000 Air Transport	11,380,568,630	675,500,000	12,056,068,630	11,380,568,630	675,500,000	12,056,068,630
	0216000 Road Safety	2,322,025,440	1,671,000,000	3,993,025,440	2,522,025,440	1,671,000,000	4,193,025,440
1093	State Department for Shipping and Maritime Affairs	2,269,974,935	1,324,000,000	3,593,974,935	2,269,974,935	1,324,000,000	3,593,974,935
	0220000 Shipping and Maritime Affairs	2,269,974,935	1,324,000,000	3,593,974,935	2,269,974,935	1,324,000,000	3,593,974,935
1094	State Department for Housing and Urban Development	1,315,392,681	86,274,800,000	87,590,192,681	1,315,392,681	86,250,800,000	87,566,192,681
	0102000 Housing Development and Human Settlement	822,678,602	76,780,300,000	77,602,978,602	822,678,602	76,816,300,000	77,638,978,602
	0105000 Urban and Metropolitan Development	146,022,769	9,494,500,000	9,640,522,769	146,022,769	9,434,500,000	9,580,522,769
	0106000 General Administration Planning and Support Services	346,691,310	-	346,691,310	346,691,310	-	346,691,310
1095	State Department for Public Works	2,069,978,552	1,205,100,000	3,275,078,552	3,699,978,552	1,309,100,000	5,009,078,552
	0103000 Government Buildings	586,054,776	586,740,000	1,172,794,776	586,054,776	586,740,000	1,172,794,776

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	0104000 Coastline Infrastructure and Pedestrian Access	92,711,281	492,260,000	584,971,281	92,711,281	596,260,000	688,971,281
	0106000 General Administration Planning and Support Services	360,590,743	4,000,000	364,590,743	360,590,743	4,000,000	364,590,743
	0218000 Regulation and Development of the Construction Industry	1,030,621,752	122,100,000	1,152,721,752	2,660,621,752	122,100,000	2,782,721,752
1104	State Department for Irrigation	1,146,882,500	18,870,780,000	20,017,662,500	1,211,382,500	22,170,780,000	23,382,162,500
	1014000 Irrigation and Land Reclamation	608,836,660	16,106,780,000	16,715,616,660	623,336,660	17,106,780,000	17,730,116,660
	015000 Water Storage and Flood Control	362,900,000	1,404,000,000	1,766,900,000	412,900,000	1,504,000,000	1,916,900,000
	1022000 Water Harvesting and Storage for Irrigation	18,224,860	1,360,000,000	1,378,224,860	18,224,860	3,560,000,000	3,578,224,860
	1023000 General Administration, Planning and Support Services	156,920,980	-	156,920,980	156,920,980	-	156,920,980
1109	State Department for Water & Sanitation	5,480,238,911	43,979,400,000	49,459,638,911	5,753,338,911	46,656,400,000	52,409,738,911
	1001000 General Administration, Planning and Support Services	538,153,125	130,000,000	668,153,125	544,153,125	130,000,000	674,153,125
	1004000 Water Resources Management	1,815,834,006	10,577,000,000	12,392,834,006	1,831,834,006	10,577,000,000	12,408,834,006
	1017000 Water and Sewerage Infrastructure Development	3,126,251,780	33,272,400,000	36,398,651,780	3,377,351,780	35,949,400,000	39,326,751,780
1112	State Department for Lands and	4,166,400,000	4,390,636,000	8,557,036,000	4,166,400,000	4,604,136,000	8,770,536,000

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	Physical Planning						
	0101000 Land Policy and Planning	2,960,044,714	3,549,376,000	6,509,420,714	2,960,044,714	3,954,376,000	6,914,420,714
	0121000 Land Information Management	-	841,260,000	841,260,000	-	649,760,000	649,760,000
	0122000 General Administration, Planning and Support Services	1,206,355,286	-	1,206,355,286	1,206,355,286	-	1,206,355,286
1122	State Department for Information Communication Technology & Digital Economy	3,355,220,752	19,182,660,000	22,537,880,752	3,525,220,752	18,829,660,000	22,354,880,752
	0207000 General Administration Planning and Support Services	278,922,194	-	278,922,194	278,922,194	-	278,922,194
	0210000 ICT Infrastructure Development	553,045,150	16,685,400,000	17,238,445,150	723,045,150	16,685,400,000	17,408,445,150
	0217000 E-Government Services	2,523,253,408	2,497,260,000	5,020,513,408	2,523,253,408	2,144,260,000	4,667,513,408
1123	State Department for Broadcasting & Telecommunications	5,179,410,364	651,900,000	5,831,310,364	5,409,410,364	651,900,000	6,061,310,364
	0207000 General Administration Planning and Support Services	231,885,786	-	231,885,786	231,885,786	-	231,885,786
	0208000 Information and Communication Services	4,731,317,827	446,900,000	5,178,217,827	4,961,317,827	446,900,000	5,408,217,827
	0209000 Mass Media Skills Development	216,206,751	205,000,000	421,206,751	216,206,751	205,000,000	421,206,751
1132	State Department for Sports	1,085,286,404	16,638,400,000	17,723,686,404	1,105,286,404	16,638,400,000	17,743,686,404

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	0901000 Sports	1,085,286,404	16,638,400,000	17,723,686,404	1,105,286,404	16,638,400,000	17,743,686,404
1134	State Department for Culture and Heritage	2,922,854,321	132,843,000	3,055,697,321	2,868,154,321	162,843,000	3,030,997,321
	0902000 Culture / Heritage	1,829,470,312	132,843,000	1,962,313,312	1,774,770,312	132,843,000	1,907,613,312
	0903000 The Arts	300,586,020	-	300,586,020	350,586,020	20,000,000	370,586,020
	0904000 Library Services	498,075,884	-	498,075,884	448,075,884	10,000,000	458,075,884
	0905000 General Administration, Planning and Support Services	166,222,722	-	166,222,722	166,222,722	-	166,222,722
	0916000 Public Records Mangement	128,499,383	-	128,499,383	128,499,383	-	128,499,383
1135	State Department for Youth Affairs and Creative Economy	1,843,510,229	2,044,961,000	3,888,471,229	1,903,510,229	2,144,961,000	4,048,471,229
	0221000 Film Development Services	500,087,720	149,731,000	649,818,720	560,087,720	149,731,000	709,818,720
	0711000 Youth Empowerment Services	339,011,526	140,160,510	479,172,036	339,011,526	240,160,510	579,172,036
	0748000 Youth Development Services	733,160,151	1,755,069,490	2,488,229,641	733,160,151	1,755,069,490	2,488,229,641
	0749000 General Administration, Planning and Support Services	271,250,832	-	271,250,832	271,250,832	-	271,250,832
1152	State Department for Energy	9,834,434,710	53,901,080,497	63,735,515,207	9,894,434,710	59,769,080,497	69,663,515,207
	0211000 General Administration Planning and Support Services	381,790,584	180,000,000	561,790,584	381,790,584	180,000,000	561,790,584
	0212000 Power Generation	1,969,736,011	13,247,750,000	15,217,486,011	2,029,736,011	13,097,750,000	15,127,486,011

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	0213000 Power Transmission and Distribution	7,414,191,396	38,338,330,497	45,752,521,893	7,414,191,396	44,416,330,497	51,830,521,893
	0214000 Alternative Energy Technologies	68,716,719	2,135,000,000	2,203,716,719	68,716,719	2,075,000,000	2,143,716,719
1162	State Department for Livestock Development	4,059,904,089	6,524,450,000	10,584,354,089	5,338,504,089	6,978,450,000	12,316,954,089
	0112000 Livestock Resources Management and Development	4,059,904,089	6,524,450,000	10,584,354,089	5,338,504,089	6,978,450,000	12,316,954,089
1166	State Department for the Blue Economy and Fisheries	2,316,495,869	9,037,930,000	11,354,425,869	2,358,795,869	9,637,930,000	11,996,725,869
	0111000 Fisheries Development and Management	2,033,407,243	8,017,420,000	10,050,827,243	2,075,707,243	8,070,420,000	10,146,127,243
	0117000 General Administration, Planning and Support Services	221,040,100	-	221,040,100	221,040,100	-	221,040,100
	0118000 Development and Coordination of the Blue Economy	62,048,526	1,020,510,000	1,082,558,526	62,048,526	1,567,510,000	1,629,558,526
1169	State Department for Crop Development	16,140,696,299	31,620,728,896	47,761,425,195	16,258,696,299	31,971,728,896	48,230,425,195
	0107000 General Administration Planning and Support Services	6,059,438,217	3,541,592,000	9,601,030,217	6,294,438,217	3,172,592,000	9,467,030,217
	0108000 Crop Development and Management	4,706,078,838	27,222,536,896	31,928,615,734	4,589,078,838	27,942,536,896	32,531,615,734
	0109000 Agribusiness and Information Management	140,139,289	747,000,000	887,139,289	140,139,289	747,000,000	887,139,289

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	0120000 Agricultural Research & Development	5,235,039,955	109,600,000	5,344,639,955	5,235,039,955	109,600,000	5,344,639,955
1173	State Department for Cooperatives	3,734,183,583	336,770,000	4,070,953,583	3,734,183,583	346,770,000	4,080,953,583
	0304000 Cooperative Development and Management	3,734,183,583	336,770,000	4,070,953,583	3,734,183,583	346,770,000	4,080,953,583
1174	State Department for Trade	3,073,661,146	-	3,073,661,146	3,098,361,146	500,000,000	3,598,361,146
	0309000 Domestic Trade and Enterprise Development	1,786,282,549	-	1,786,282,549	1,786,282,549	-	1,786,282,549
	0310000 Fair Trade Practices And Compliance of Standards	116,359,164	-	116,359,164	116,359,164	-	116,359,164
	0311000 International Trade Development and Promotion	802,937,016	-	802,937,016	827,637,016	500,000,000	1,327,637,016
	0312000 General Administration, Planning and Support Services	368,082,417	-	368,082,417	368,082,417	-	368,082,417
1175	State Department for Industry	2,002,606,621	6,555,690,000	8,558,296,621	2,072,906,621	6,366,770,000	8,439,676,621
	0301000 General Administration Planning and Support Services	377,045,857	-	377,045,857	377,045,857	-	377,045,857
	0320000 Industrial Promotion and Development	751,543,169	4,811,880,000	5,563,423,169	793,743,169	4,822,960,000	5,616,703,169
	0321000 Standards and Quality Infrastructure & Research	874,017,595	1,743,810,000	2,617,827,595	902,117,595	1,543,810,000	2,445,927,595
1176	State Department for Micro, Small	1,529,618,500	7,696,840,000	9,226,458,500	1,566,218,500	7,802,840,000	9,369,058,500

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	and Medium Enterprises Development						
	0316000 Promotion and Development of MSMEs	493,739,288	2,516,840,000	3,010,579,288	498,739,288	2,522,840,000	3,021,579,288
	0317000 Product and Market Development for MSMEs	355,726,000	80,000,000	435,726,000	387,326,000	80,000,000	467,326,000
	0318000 Digitization and Financial Inclusion for MSMEs	350,700,000	5,100,000,000	5,450,700,000	350,700,000	5,200,000,000	5,550,700,000
	0319000 General Administration, Planning and Support Services	329,453,212	-	329,453,212	329,453,212	-	329,453,212
1177	State Department for Investment Promotion	960,613,914	3,105,430,000	4,066,043,914	1,010,613,914	3,605,430,000	4,616,043,914
	0322000 Investment Development and Promotion	960,613,914	3,105,430,000	4,066,043,914	1,010,613,914	3,605,430,000	4,616,043,914
1184	State Department for Labour and Skills Development	4,219,529,843	1,512,885,400	5,732,415,243	4,319,529,843	1,512,885,400	5,832,415,243
	0910000 General Administration Planning and Support Services	509,933,940	-	509,933,940	509,933,940	-	509,933,940
	0906000 Labour, Employment and Safety Services	1,069,422,931	346,105,400	1,415,528,331	1,169,422,931	346,105,400	1,515,528,331
	0907000 Manpower Development, Industrial Skills & Productivity Management	2,640,172,972	1,166,780,000	3,806,952,972	2,640,172,972	1,166,780,000	3,806,952,972
1185	State Department for Social Protection and	33,064,925,645	1,939,880,000	35,004,805,645	33,114,925,645	2,189,880,000	35,304,805,645

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	Senior Citizen Affairs						
	0908000 Social Development and Children Services	4,432,041,070	32,259,000	4,464,300,070	4,482,041,070	282,259,000	4,764,300,070
	0909000 National Social Safety Net	28,381,293,750	1,907,621,000	30,288,914,750	28,381,293,750	1,907,621,000	30,288,914,750
	0914000 General Administration, Planning and Support Services	251,590,825	-	251,590,825	251,590,825	-	251,590,825
1192	State Department for Mining	905,898,447	652,260,000	1,558,158,447	1,105,898,447	652,260,000	1,758,158,447
	1007000 General Administration Planning and Support Services	356,382,218	-	356,382,218	436,382,218	-	436,382,218
	1009000 Mineral Resources Management	266,659,867	270,800,000	537,459,867	331,659,867	270,800,000	602,459,867
	1021000 Geological Survey and Geoinformation Management	282,856,362	381,460,000	664,316,362	337,856,362	381,460,000	719,316,362
1193	State Department for Petroleum	27,325,211,883	3,875,200,000	31,200,411,883	27,325,211,883	3,875,200,000	31,200,411,883
	0215000 Exploration and Distribution of Oil and Gas	27,325,211,883	3,875,200,000	31,200,411,883	27,325,211,883	3,875,200,000	31,200,411,883
1202	State Department for Tourism	9,858,821,808	470,000,000	10,328,821,808	9,858,821,808	470,000,000	10,328,821,808
	0313000 Tourism Promotion and Marketing	536,260,000	30,000,000	566,260,000	536,260,000	30,000,000	566,260,000
	0314000 Tourism Product Development and Diversification	9,053,712,966	440,000,000	9,493,712,966	9,053,712,966	440,000,000	9,493,712,966

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	0315000 General Administration, Planning and Support Services	268,848,842	-	268,848,842	268,848,842	-	268,848,842
1203	State Department for Wildlife	12,209,194,935	1,333,000,000	13,542,194,935	12,090,194,935	2,253,000,000	14,343,194,935
	1019000 Wildlife Conservation and Management	12,209,194,935	1,333,000,000	13,542,194,935	12,090,194,935	2,253,000,000	14,343,194,935
1212	State Department for Gender and Affirmative Action	1,940,841,404	3,518,700,000	5,459,541,404	2,075,841,404	4,038,700,000	6,114,541,404
	0911000 Community Development	43,400,000	3,000,000,000	3,043,400,000	983,400,000	3,500,000,000	4,483,400,000
	0912000 Gender Empowerment	1,643,486,225	518,700,000	2,162,186,225	838,486,225	538,700,000	1,377,186,225
	0913000 General Administration, Planning and Support Services	253,955,179	-	253,955,179	253,955,179	-	253,955,179
1213	State Department for Public Service	17,076,244,125	1,138,445,784	18,214,689,909	18,371,244,125	1,241,445,784	19,612,689,909
	0710000 Public Service Transformation	6,596,629,218	1,019,645,784	7,616,275,002	7,646,629,218	1,122,645,784	8,769,275,002
	0709000 General Administration Planning and Support Services	384,125,877	60,000,000	444,125,877	384,125,877	60,000,000	444,125,877
	0747000 National Youth Service	10,095,489,030	58,800,000	10,154,289,030	10,340,489,030	58,800,000	10,399,289,030
1221	State Department for East African Community	612,087,899	35,400,000	647,487,899	612,087,899	35,400,000	647,487,899
	0305000 East African Affairs and Regional Integration	612,087,899	35,400,000	647,487,899	612,087,899	35,400,000	647,487,899
1252	State Law Office	5,320,570,997	157,000,000	5,477,570,997	6,820,570,997	157,000,000	6,977,570,997

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	0606000 Legal Services	2,968,584,833	-	2,968,584,833	4,468,584,833	-	4,468,584,833
	0607000 Governance, Legal Training and Constitutional Affairs	1,360,207,486	35,000,000	1,395,207,486	1,360,207,486	35,000,000	1,395,207,486
	0609000 General Administration, Planning and Support Services	991,778,678	122,000,000	1,113,778,678	991,778,678	122,000,000	1,113,778,678
1271	Ethics and Anti-Corruption Commission	3,913,630,000	57,920,000	3,971,550,000	4,113,630,000	57,920,000	4,171,550,000
	0611000 Ethics and Anti-Corruption	3,913,630,000	57,920,000	3,971,550,000	4,113,630,000	57,920,000	4,171,550,000
1281	National Intelligence Service	45,851,000,000	-	45,851,000,000	46,351,000,000	-	46,351,000,000
	0804000 National Security Intelligence	45,851,000,000	-	45,851,000,000	46,351,000,000	-	46,351,000,000
1291	Office of the Director of Public Prosecutions	3,859,020,000	48,500,000	3,907,520,000	3,959,020,000	48,500,000	4,007,520,000
	0612000 Public Prosecution Services	3,859,020,000	48,500,000	3,907,520,000	3,959,020,000	48,500,000	4,007,520,000
1311	Office of the Registrar of Political Parties	1,037,871,453	-	1,037,871,453	2,037,871,453	-	2,037,871,453
	0614000 Registration, Regulation and Funding of Political Parties	1,037,871,453	-	1,037,871,453	2,037,871,453	-	2,037,871,453
1321	Witness Protection Agency	741,192,500	-	741,192,500	741,192,500	-	741,192,500
	0615000 Witness Protection	741,192,500	-	741,192,500	741,192,500	-	741,192,500
1331	State Department for Environment &	3,082,335,109	1,686,796,186	4,769,131,295	3,332,335,109	1,686,796,186	5,019,131,295

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	Climate Change						
	1002000 Environment Management and Protection	1,547,857,770	1,170,296,186	2,718,153,956	1,547,857,770	1,170,296,186	2,718,153,956
	1010000 General Administration, Planning and Support Services	436,386,346	-	436,386,346	686,386,346	-	686,386,346
	1012000 Meteorological Services	1,098,090,993	486,000,000	1,584,090,993	1,098,090,993	486,000,000	1,584,090,993
	1018000 Forests Development, Management and Conservation	-	30,500,000	30,500,000	-	30,500,000	30,500,000
1332	State Department for Forestry	9,043,630,000	3,520,300,000	12,563,930,000	9,043,630,000	3,520,300,000	12,563,930,000
	1018000 Forests and Water Towers Conservation	8,896,148,407	3,520,300,000	12,416,448,407	8,896,148,407	3,520,300,000	12,416,448,407
	1024000 Agroforestry and Commercial Forestry Development	11,119,622	-	11,119,622	11,119,622	-	11,119,622
	1025000 General Administration, Planning and Support Services	136,361,971	-	136,361,971	136,361,971	-	136,361,971
2011	Kenya National Commission on Human Rights	478,074,025	-	478,074,025	478,074,025	-	478,074,025
	0616000 Protection and Promotion of Human Rights	478,074,025	-	478,074,025	478,074,025	-	478,074,025
2021	National Land Commission	1,506,262,679	103,460,000	1,609,722,679	1,868,362,679	147,860,000	2,016,222,679
	0119000 Land Administration and Management	1,506,262,679	103,460,000	1,609,722,679	1,868,362,679	147,860,000	2,016,222,679
2031	Independent Electoral and	3,730,899,680	24,320,000	3,755,219,680	3,730,899,680	24,320,000	3,755,219,680

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	Boundaries Commission						
	0617000 Management of Electoral Processes	3,694,852,948	24,320,000	3,719,172,948	3,694,852,948	24,320,000	3,719,172,948
	0618000 Delimitation of Electoral Boundaries	36,046,732	-	36,046,732	36,046,732	-	36,046,732
2061	The Commission on Revenue Allocation	413,465,304	-	413,465,304	413,465,304	-	413,465,304
	0737000 Inter-Governmental Transfers and Financial Matters	413,465,304	-	413,465,304	413,465,304	-	413,465,304
2071	Public Service Commission	3,622,230,017	45,300,000	3,667,530,017	3,622,230,017	45,300,000	3,667,530,017
	0725000 General Administration, Planning and Support Services	855,023,567	45,300,000	900,323,567	855,023,567	45,300,000	900,323,567
	0726000 Human Resource Management and Development	2,492,690,989	-	2,492,690,989	2,492,690,989	-	2,492,690,989
	0727000 Governance and National Values	171,923,179	-	171,923,179	171,923,179	-	171,923,179
	0744000 Performance and Productivity Management	56,512,561	-	56,512,561	56,512,561	-	56,512,561
	075000 Administration of Quasi-Judicial Functions	46,079,721	-	46,079,721	46,079,721	-	46,079,721
2081	Salaries and Remuneration Commission	472,230,922	-	472,230,922	472,230,922	-	472,230,922
	0728000 Salaries and Remuneration Management	472,230,922	-	472,230,922	472,230,922	-	472,230,922

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
2091	Teachers Service Commission	352,473,737,118	472,329,000	352,946,066,118	357,773,737,118	442,329,000	358,216,066,118
	0509000 Teacher Resource Management	342,281,134,808	404,329,000	342,685,463,808	347,281,134,808	404,329,000	347,685,463,808
	0510000 Governance and Standards	1,309,066,143	-	1,309,066,143	1,309,066,143	-	1,309,066,143
	0511000 General Administration, Planning and Support Services	8,883,536,167	68,000,000	8,951,536,167	9,183,536,167	38,000,000	9,221,536,167
2101	National Police Service Commission	1,131,272,317	-	1,131,272,317	1,131,272,317	-	1,131,272,317
	0620000 National Police Service Human Resource Management	1,131,272,317	-	1,131,272,317	1,131,272,317	-	1,131,272,317
2111	Auditor General	8,211,770,850	355,000,000	8,566,770,850	8,211,770,850	455,000,000	8,666,770,850
	0729000 Audit Services	8,211,770,850	355,000,000	8,566,770,850	8,211,770,850	455,000,000	8,666,770,850
2121	Office of the Controller of Budget	652,219,080	-	652,219,080	740,219,080	-	740,219,080
	0730000 Control and Management of Public finances	652,219,080	-	652,219,080	740,219,080	-	740,219,080
2131	Commission on Administrative Justice	661,974,500	-	661,974,500	661,974,500	-	661,974,500
	0731000 Promotion of Administrative Justice	661,974,500	-	661,974,500	661,974,500	-	661,974,500
2141	National Gender and Equality Commission	425,810,000	10,000,000	435,810,000	425,810,000	10,000,000	435,810,000
	0621000 Promotion of Gender Equality and Freedom	425,810,000	10,000,000	435,810,000	425,810,000	10,000,000	435,810,000

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	from Discrimination						
2151	Independent Policing Oversight Authority	1,107,672,060	-	1,107,672,060	1,107,672,060	-	1,107,672,060
	0622000 Policing Oversight Services	1,107,672,060	-	1,107,672,060	1,107,672,060	-	1,107,672,060
	Sub-Total: Executive	1,518,676,252,608	724,378,630,302	2,243,054,882,910	1,579,183,298,315	744,410,910,302	2,323,594,208,617
1261	The Judiciary	21,287,400,000	1,500,000,000	22,787,400,000	22,137,400,000	1,600,000,000	23,737,400,000
	0610000 Dispensation of Justice	21,287,400,000	1,500,000,000	22,787,400,000	22,137,400,000	1,600,000,000	23,737,400,000
2051	Judicial Service Commission	902,900,000	-	902,900,000	902,900,000	-	902,900,000
	0619000 General Administration, Planning and Support Services	902,900,000	-	902,900,000	902,900,000	-	902,900,000
	Sub-Total: Judiciary	22,190,300,000	1,500,000,000	23,690,300,000	23,040,300,000	1,600,000,000	24,640,300,000
2041	Parliamentary Service Commission	3,062,145,629	-	3,062,145,629	1,167,000,000	-	1,167,000,000
	0765000 General Administration, Planning and Support Services	2,806,577,429	-	2,806,577,429	1,127,000,000	-	1,127,000,000
	0766000 Human Resource Management and Development	255,568,200	-	255,568,200	40,000,000	-	40,000,000
2042	National Assembly	32,486,292,127	-	32,486,292,127	26,775,000,000	-	26,775,000,000
	0721000 National Legislation, Representation and Oversight	32,486,292,127	-	32,486,292,127	26,775,000,000	-	26,775,000,000
2043	Parliamentary Joint Services	11,058,000,000	4,785,000,000	15,843,000,000	6,581,000,000	2,065,000,000	8,646,000,000
	0723000 General Administration,	10,650,117,052	4,785,000,000	15,435,117,052	6,363,811,050	2,065,000,000	8,428,811,050

SCHEDULE: BUDGET ESTIMATES FOR FY 2024/25

VOTE CODE	VOTE & PROGRAMME	SUBMITTED BUDGET ESTIMATES FOR FY 2024/25			FINAL BUDGET ESTIMATES FOR FY 2024/25		
		CURRENT	CAPITAL	TOTAL	CURRENT	CAPITAL	TOTAL
	Planning and Support Services						
	0746000 Legislative Training Research & Knowledge Management	407,882,948	-	407,882,948	217,188,950	-	217,188,950
2044	Senate	14,418,972,002	-	14,418,972,002	8,010,000,000	-	8,010,000,000
	0767000 Senate Legislation and Oversight	6,123,327,497	-	6,123,327,497	3,236,650,000	-	3,236,650,000
	0768000 Senate Representation, Liaison and Intergovernmental Relations	3,512,131,036	-	3,512,131,036	2,178,199,100	-	2,178,199,100
	0769000 General Administration, Planning and Support Services	4,783,513,469	-	4,783,513,469	2,595,150,900	-	2,595,150,900
	Sub-Total: Parliament	61,025,409,758	4,785,000,000	65,810,409,758	42,533,000,000	2,065,000,000	44,598,000,000
Grand Total		1,601,891,962,366	730,663,630,302	2,332,555,592,668	1,644,756,598,315	748,075,910,302	2,392,832,508,617

NOTICES

I. THE KENYA DRUGS AUTHORITY BILL (NATIONAL ASSEMBLY BILL NO. 54 OF 2022)

- 1) Notice is given that the Chairperson of the Departmental Committee on Health intends to move the following amendment to the Kenya Drugs Authority Bill, 2022 at the Committee Stage—

LONG TITLE

THAT, the Bill be amended by deleting the Long Title and substituting therefor the following new Long Title—

“AN ACT of Parliament to establish a comprehensive legal framework for the regulation of health products and technologies; to safeguard public health through development of a regulatory system to ensure safety, quality, efficacy, effectiveness and performance of health products; to establish the Kenya Health Products and Technologies Regulatory Authority and for connected purposes”.

CLAUSE 1

THAT, Clause 1 of the Bill be amended by—

- (a) deleting the phrase “Kenya Drugs Authority Act, 2022” and substituting therefor the phrase “Kenya Health Products and Technologies Regulatory Authority Act, 2022”;
- (b) deleting the words “and commencement” in the marginal note.

CLAUSE 2

THAT, Clause 2 of the Bill be amended—

- (a) in the definition of “article” by—
 - (i) inserting the words “dietary supplement” immediately after the words “therapeutic cosmetic” appearing in paragraph (a); and
 - (ii) inserting the words “dietary supplement” immediately after the words “therapeutic cosmetic” appearing in paragraph (b);
- (b) in the definition of “Authority” by deleting the words “Kenya Drugs Authority” and substituting therefor the words, “Kenya Health Products and Technologies Regulatory Authority”;
- (c) in the definition of “chemical substance” by deleting the words “or detergent”;
- (d) in the definition of “drug” by deleting the word “if” appearing in paragraph (b)(ii) and substituting therefor the word “of”;

- (e) by deleting the definition of “enrolled pharmaceutical technologist”;
- (f) in the definition of “health products and technologies” by inserting the words, “dietary supplements” immediately after the words, “therapeutic cosmetics”;
- (g) by deleting the definition of “herbal medicine or product”;
- (h) by deleting the definition of “medical device”;
- (i) by deleting the definition of “medicinal substance”;
- (j) in the definition of “package” by inserting the words “dietary supplement” immediately after the words “therapeutic cosmetic”;
- (k) by deleting the definition of “pharmacy”;
- (l) by deleting the definition of “pharmaceutical technologist”;
- (m) by deleting the definition of “registered midwife”;
- (n) in the definition of “scheduled substance” by deleting the phrase “in the relevant schedule under this Act” and substituting therefor the phrase “in the list published by the Cabinet Secretary under section 37”;
- (o) by deleting the definition of “therapeutic cosmetic”; and
- (p) by inserting the following new definitions in their proper alphabetical sequence—

“active surveillance” means prospective measures taken to detect adverse drug reactions and adverse events and involves active follow-up during and after treatment of patients where the events may be detected by asking the patient directly or screening patient records;

“adverse drug reaction” means a response to a drug which is noxious and unintended, and which occurs at doses normally used in humans for the prophylaxis, diagnosis or therapy of disease, or for the modification of physiological function and is characterized by the suspicion of a causal relationship between a medical product and an occurrence;

“adverse event” means any untoward medical occurrence that may present during treatment with a pharmaceutical product but which does not necessarily have a causal relationship with the treatment;

“alternative medicine” means complementary medicine and includes a broad set of health care practices that are not part of Kenya’s tradition and are not integrated into dominant health care system;

“biologicals” means a diverse group of medicines which includes vaccines, growth factors, immune modulators, monoclonal antibodies and includes products derived from human blood and plasma;

“Board” means the Board of the Authority established under section 8;

“Centre” means the National Pharmacovigilance Centre established under section 59B;

“clinical trial” means any systematic study on pharmaceutical products in human subjects, whether in patients or other volunteers, in order to discover or verify the effects of, identify any adverse reaction to investigational products, study the absorption, distribution, metabolism and excretion of the products with the object of ascertaining their efficacy and safety;

“dietary supplement” means a product taken by mouth that is added to the diet to help meet daily requirements of essential nutrients, and which usually contains one or more dietary ingredient and includes vitamins, minerals and herbs;

“enrolled pharmaceutical technologist” means a person enrolled as such by the body for the time being responsible for the enrolment of pharmaceutical technologists;

“falsified medical product” means a product that is deliberately or fraudulently misrepresented in relation to its identity, composition or source;

“Field Safety Corrective Action” means any action taken by a product owner to reduce a risk of death or serious deterioration in the state of health associated with the use of a medical device, and includes—

- (a) the return of a medical device to the product owner or its representative;
- (b) device modification which may include—
 - (i) retrofit in accordance with the product owner’s modification or design change;
 - (ii) permanent or temporary changes to the labelling or instructions for use;
 - (iii) software upgrades including those carried out by remote access;
 - (iv) modification to the clinical management of patients to address a risk of serious injury or death related specifically to the characteristics of the device;
 - (v) device exchange;
 - (vi) device destruction; or

- (vii) advice given by product owner regarding the use of the device.

“health product” includes a medicine, medical product, medicinal substance, vaccine, diagnostic, medical device, blood or blood product, herbal medicine, therapeutic feed and nutritional formulation, cosmetic and related products;

“health technology” means the application of organized knowledge and skills in the form of medicines, devices, vaccines, procedures, and systems developed to solve a health problem and improve the quality of lives, and includes radiation-emitting devices and related products;

“herbal medicine or product” means a plant derived material or preparations with claimed therapeutic or other health benefits, which contain either raw or processed ingredients from one or more plants or material of inorganic or animal origin and includes herbs, herbal materials, herbal preparations, finished herbal products that contain active ingredients, parts of plants or other plant materials or combinations;

“Inspector of Drugs” means a person who is competitively recruited by the Authority as a drug inspector under this Act;

“lot” or “sub-lot” means a defined quantity of starting material, packaging material or product, processed in a single process or series of processes so that the quantity is expected to be homogeneous; and in the case of continuous manufacture, the lot corresponds to a defined fraction of the production characterized by its intended homogeneity;

“lot release” means the process of the evaluation of an individual lot of a licensed biological product by the Authority before giving approval for its release onto the market;

“marketing authorization” means the certificate of registration issued by the competent health product regulatory authority in the country of origin for the purpose of marketing or free distribution of a health product after evaluation for safety, efficacy and quality;

“medical device” means any instrument, apparatus, implement, machine, appliance, implant, reagent for in vitro use, software, material or other similar or related article, intended by the manufacturer to be used, alone or in combination, for human beings, for one or more of the specific medical purpose of—

- (a) diagnosis, prevention, monitoring, treatment or alleviation of disease;
- (b) diagnosis, monitoring, treatment, alleviation of or compensation for an injury;
- (c) investigation, replacement, modification or support of the anatomy or of a physiological process;
- (d) supporting or sustaining life;
- (e) control of conception;
- (f) disinfection of medical devices;
- (g) providing information by means of in vitro examination of specimens derived from the human body;
- (h) disinfection substances;
- (i) aids for persons with disabilities;
- (j) devices incorporating animal or human tissues;
- (k) devices for in-vitro fertilization or assisted reproduction technologies,

and does not achieve its primary intended action by pharmacological, immunological or metabolic means, in or on the human body, but which may be assisted in its intended function by such means;

“medicinal substance” means a substance, the origin of which may be human, animal, vegetable or chemical including human blood and human blood products, micro-organisms, whole animals, parts of organs, animal secretions, toxins, extracts, blood products, micro-organisms, plants, parts of plants, vegetable secretions, extracts, elements, naturally occurring chemical materials and chemical products obtained by chemical change or synthesis;

“passive surveillance” means that no active measures are taken to look for adverse effects other than the encouragement of health professionals and others to report safety concerns;

“parallel importation” means importation into Kenya, by a licensed importer of a health product other than the marketing authorization holder or his or her technical representative, of the following health products which require marketing authorization in Kenya—

- (a) patented health products under the applicable law;
- (b) non-patented health products; or
- (c) branded generic health products;

“parallel imported medicinal substance” means a medicinal substance imported into Kenya under this Act;

“pharmacovigilance” means the science and activities relating to the detection, assessment, understanding and prevention of adverse effects or any other possible health product related problem;

“premise” includes any land, building, dwelling-place or any other place whatsoever; and includes stand-alone community retail pharmacy, private hospital pharmacy, public health facility pharmacy, wholesale pharmacy or distribution outlet, where health products and technologies are stored, handled or distributed;

“Registrar” means the Director-General of the Authority appointed under section 6;

“scheduling” means, in relation to a substance, the determination of the schedule or schedules to the current Poisons Standard in which the name or a description of the substance is to be included;

“therapeutic cosmetic” means a product with the ability to trigger biological actions on the dermis, skin, eyes or teeth, to prevent future damage and contains ingredients that are usually not found in regular cosmetics or at higher strengths than could be sold safely over the counter;

“traditional medicine” includes the knowledge, skills and practices based on the theories, beliefs and experiences indigenous to different cultures, whether explicable or not, used in the maintenance of health as well as in the prevention, diagnosis, improvement or treatment of physical and mental illness;

“unregistered medical product” means a product that has not undergone evaluation and approval by the Authority subject to permitted conditions under the Act and the rules therein;

“vessel” means a truck, van, bus, minibus, car, trailer, aircraft, railway carriage, boat and other means that are used for purposes of conveying health products and technologies;

“wholesale dealer” means a person who is licensed to carry out a business where health products and technologies are stored, distributed or sold in bulk to persons other than individual consumers and includes registration, importation, warehousing, good distribution practices and pharmacovigilance;”.

CLAUSE 3

THAT, Clause 3 of the Bill be amended—

(a) by deleting sub-clause (1) and substituting therefor the following new sub-clause (1)—

“(1) This Act applies to the regulation of—

- (a) medical devices including radiation emitting devices;
- (b) radiopharmaceuticals;
- (c) complementary or herbal medicines;
- (d) cosmetics and borderline products;
- (e) in-vitro diagnostics medical devices;
- (f) therapeutic feeds;
- (g) clinical trials;
- (h) nutraceuticals and dietary supplements;
- (i) digital health and technologies;
- (j) scheduled substances;
- (k) chemical substances; and
- (l) biological products for use in humans and the starting materials used in their manufacture.”

(b) by inserting the following new sub-clause immediately after sub-clause (2)—

“(3) This Act shall not apply to the regulation of traditional medicine and alternative medicine.”

CLAUSE 4

THAT, Clause 4 of the Bill be amended in sub-clause (1) by deleting the words “Kenya Drugs Authority” and substituting therefor the words “Kenya Health Products and Technologies Regulatory Authority”.

CLAUSE 5

THAT, Clause 5 of the Bill be amended by deleting the words, “but the Authority may establish branches anywhere in Kenya” and substituting therefor the words “or in such other place as the Board of the Authority may, by resolution, determine”.

CLAUSE 6

THAT, Clause 6 of the Bill be amended—

- (a) by deleting sub-clause (1) and substituting therefor the following new sub-clause (1)—

“(1) There shall be a Director-General of the Authority who shall be the chief executive officer of the Authority.”

- (b) by deleting sub-clause (2) and substituting therefor the following new sub-clause (2)—

“(2) The Director-General shall be appointed by the Board, through a transparent and competitive process, on such terms as may be specified in the instrument of appointment.”

(c) in sub-clause (3) by deleting the word “four” and substituting therefor the word “three”.

(d) by deleting sub-clause (4) and substituting the following new sub-clause (4)—

“(4) A person shall be qualified for appointment as a Director-General if such person—

- (a) holds a bachelor’s degree in pharmacy from a university recognized in Kenya;
- (b) holds a masters’ degree in pharmacy, medicine or any relevant field from a university recognized in Kenya;
- (c) has at least ten years’ experience in pharmacy or its equivalent;
- (d) has served in a senior management position for at least five years;
- (e) is a member of a professional body; and
- (f) meets the requirements of Chapter six of the Constitution.”; and

(e) by deleting sub-clause (5).

CLAUSE 7

THAT, Clause 7 of the Bill be amended in paragraph (f) by deleting the phrase “Act. regulation under this” and substituting therefor the phrase “regulation under this Act.”

CLAUSE 8

THAT, Clause 8 of the Bill be amended—

(a) by deleting sub-clause (1) and substituting therefor the following new sub-clause—

“(1) The management of the Authority shall vest in a Board appointed under this section.”

(b) by deleting sub-clause (2) and substituting therefor the following new sub-clause (2)—

“(2) The Board shall comprise—

- (a) a non-executive Chairperson appointed by the President and who shall—

- (i) be a registered pharmacist of good standing with a degree in pharmacy; and
 - (ii) have at least ten years' experience in the pharmaceutical sector, five of which shall be at senior management level;
 - (b) the Principal Secretary in the Ministry for the time being responsible for health or a representative designated in writing;
 - (c) the Principal Secretary in the Ministry for the time being responsible for finance or a representative designated in writing;
 - (d) the Director-General for Health or a representative designated in writing;
 - (e) one person nominated by the Pharmaceutical Society of Kenya;
 - (f) one person nominated by the Kenya Pharmaceutical Association;
 - (g) one person nominated by the Kenya Medical Association;
 - (h) one person, not being a Governor, with knowledge and experience in health products and technologies nominated by the Council of County Governors to represent the interests of counties;
 - (i) one person, not being a public officer, representing consumer protection nominated by the Consumer Federation of Kenya; and
 - (j) the Director-General of the Authority who shall be the secretary and an *ex officio* member of the Board.”; and
- (c) by deleting sub-clause (3) and substituting therefor the following new sub-clause (3)—

“(3) The Cabinet Secretary shall appoint the members of the Board under subsection (2) (e), (f), (g), (h) and (i) by notice in the *Gazette*.”

CLAUSE 9

THAT, the Bill be amended by deleting Clause 9.

CLAUSE 10

THAT, Clause 10 of the Bill be amended in sub-clause (1) by deleting the words “section 12” appearing in paragraph (c) and substituting therefor the words “section 11”.

CLAUSE 12

THAT, Clause 12 of the Bill be amended by—

- (a) inserting the following paragraphs immediately after paragraph (e)—

- “(ea) regulate the disposal of health products and technologies;
(eb) monitor the market for the presence of unregistered and illegal health products and technologies;
(ec) conduct analytical tests of health products and technologies”;
- (b) deleting paragraph (f) and substituting therefor the following new paragraph (f)
—
“(f) ensure continuous monitoring of the safety of health products and technologies regulated under this Act through analysis of reports on adverse reactions and events, including any other health product and technology use related issues and take appropriate regulatory actions when necessary”;
- (c) deleting paragraph (g) and substituting therefor the following new paragraph (g)—
“(g) regulate clinical trials and ensure that clinical trial protocols of health products and technologies are being assessed according to the prescribed ethical and professional criteria and defined standards including mandatory bioequivalence studies”;
- (d) inserting the following new paragraphs immediately after paragraph (g)—
“(ga) approve the use of any unregistered medicinal substance for purposes of clinical trials, emergency use and compassionate use;
(gb) carry out pharmacovigilance audits and inspections in order to ensure compliance with good pharmacovigilance practices and the prescribed requirements”;
- (e) deleting paragraph (n) and substituting therefor the following new paragraph (n)—
“(n) appoint inspectors who hold a minimum of a diploma in pharmacy and conduct inspection, either by itself or through its agents, of manufacturing premises, medical devices establishments, importing and exporting agents, wholesalers, distributors, pharmacies, including those in health facilities and clinics, retail outlets and any other premises and vessels subject to regulation under this Act”;
- (f) inserting the following new paragraphs after paragraph (o)—

“(oa) conduct national regulatory authority lot release, official authority batch release of specified biologicals to ensure the quality, safety and efficacy of biological products through a regulatory release system in compliance with established approaches, policies, guidelines, procedures and in line with World Health Organization and internationally recognized guidelines;
(ob) carry out and promote research related to medicines and health products”;

(g) inserting the following paragraphs after paragraph (q)—

“(qa) ensure that all health products and technologies manufactured in, imported into or exported from the country including through parallel importation conform to prescribed standards of quality, safety and efficacy;

(qb) enforce the prescribed standards of quality, safety and efficacy of health products and technologies manufactured, imported into or exported out of the country;

(qc) grant or revoke licenses and permits for the manufacture, importation, exportation, distribution and sale of health products and technologies;

(qd) maintain a register of all authorized health products and technologies manually or electronically;

(qe) regulate licit use of narcotic, psychotropic substances and precursor chemical substances in accordance with the Single Convention on Narcotic Drugs, 1961, the Convention on Psychotropic substances, 1971 or the United Nations Convention against Illicit Traffic of Precursor Chemical Substances, 1988;

(qf) inspect and license all manufacturing premises, importing and exporting agents, wholesalers, distributors, pharmacies including those in hospitals and clinics and other retail outlets;”

CLAUSE 13

THAT, Clause 13 of the Bill be amended by—

(a) deleting paragraph (a) and substituting therefor the following new paragraph (a)—

“(a) collaborate with such other bodies or organizations within or outside Kenya as it may consider desirable or appropriate for the furtherance of the purpose of the Act;”

(b) inserting the following new paragraphs immediately after paragraph (a)—

“(aa) adopt and implement any such internationally recognized good regulatory practices;
(ab) determine and implement effective and efficient reliance mechanisms;
(ac) issue, suspend, withdraw or revoke any license or compliance certificate granted under this Act;
(ad) levy, collect and utilize fees for services rendered;
(ae) grant or withdraw licenses and permits to manufacturers, wholesalers, retailers, importers, exporters and distributors;
(af) develop guidelines on the manufacture, import and export, distribution, sale and use of medical products”.

CLAUSE 21

THAT, Clause 21 of the Bill be amended—

- (a) by deleting sub-clause (1) and substituting therefor the following new sub-clause (1)—

“(1) The Board may establish such scientific advisory committees of the Authority, as may be necessary for the effective performance of the functions of the Authority”.

- (b) in sub-clause (3) by deleting the words “Cabinet Secretary” and substituting therefor the words “Board of the Authority”;
(c) in sub-clause (4) by deleting the words “Cabinet Secretary” and substituting therefor the words “Board of the Authority”;
(d) by deleting sub-clause (9) and substituting therefor the following new sub-clause (9) —
“(9) A scientific advisory committee shall submit, at least once every six months, a report to the Board of the Authority, with respect to its activities and the Board shall submit a copy of each report to the Cabinet Secretary”.

PART IV

THAT, Part IV of the Bill be amended by deleting the title and substituting therefor the following new title—

“PART III—HEALTH PRODUCTS AND TECHNOLOGIES”

CLAUSE 22

THAT, Clause 22 of the Bill be amended—

- (a) in the marginal note by deleting the word “medicines” and substituting therefor the words “health products and technologies”;

(b) in sub-clause (1) by—

- (i) deleting the words “sell any medicine” appearing in the opening sentence and substituting therefor the words “sell, manufacture, supply, distribute or dispense any health product or technology”;
- (ii) deleting paragraph (d) and substituting therefore the following new paragraph (d)—

“(d) is falsified,”;

(c) in sub-clause (3) by—

- (i) deleting the word “medicine” appearing in the opening sentence and substituting therefor the words “health product or technology”; and
- (ii) deleting the words “pharmaceutical product” appearing in paragraph (b) and substituting therefor the words, “health product or technology”.

CLAUSE 23

THAT, Clause 23 of the Bill be amended—

(a) in sub-clause (1) by—

- (i) deleting the word “medicines” appearing in paragraph (a) and substituting therefor the words, “health products or technologies”;
- (ii) deleting the word “medicine” appearing in paragraph (b) and substituting therefor the words, “health product or technology”; and
- (iii) deleting the word “medicine” appearing in paragraph (c) and substituting therefor the words, “health product or technology”;

(b) in sub-clause (2) by—

- (i) deleting the words “one million” appearing in paragraph (a) and substituting therefor the words “two million”; and
- (ii) deleting the words “two million” appearing in paragraph (b) and substituting therefor the words “five million”.

CLAUSE 24

THAT, Clause 24 of the Bill be amended—

- (a) in the marginal note by deleting the word “medicines” and substituting therefor the words “health products and technologies”;

- (b) in sub-clause (1) by deleting the word “medicine” wherever it appears and substituting therefor the words “health product or technology”;
- (c) by deleting sub-clause (2) and substituting therefor the following new sub-clause (2)—
“(2) If a standard has not been prescribed for a health product or technology but a standard for the health product or technology is contained in any of the publications specified in the Fifth Schedule, any person who manufactures, labels, packages, sells or advertises any other substance or article in such a manner that is likely to be mistaken for the health product or technology having met any of the standards contained in any of the publications specified in the Fifth Schedule, commits an offence.”;
- (d) in sub-clause (3) by—
(i) deleting the word “medicine” wherever it appears in the opening sentence and substituting therefor the words “health product or technology”; and
(ii) deleting the word “drug” appearing in paragraph (b) and substituting therefor the words “health product or technology”;
- (e) in sub-clause (4) by—
(i) deleting the phrase “one hundred thousand shillings or to imprisonment for a term not exceeding three months” appearing in paragraph (a) and substituting therefor the phrase “one million shillings or to imprisonment for a term not exceeding three years”; and
(ii) deleting the words “two hundred thousand” appearing in paragraph (b) and substituting therefor the words “two million”.

CLAUSE 25

THAT, the Bill be amended by deleting Clause 25.

CLAUSE 26

THAT, Clause 26 of the Bill be amended by—

- (a) deleting the word “medicine” appearing in the marginal note and substituting therefor the words “health product or technology”; and
(b) deleting the word “medicine” and substituting therefor the words “health product or technology”.

CLAUSE 27

THAT, Clause 27 of the Bill be amended by—

- (a) deleting the words “medicinal products” appearing in paragraph (a) and substituting therefor the words “health products or technologies”;
- (b) deleting the words “medicinal products” appearing in paragraph (b) and substituting therefor the words “health products or technologies”; and
- (c) deleting paragraph (c) and substituting therefor the following new paragraph (c)—

“(c) the quality of the health products or technologies of such description, according to the specification and the method or proposed method of manufacture of the health products or technologies, and the provisions proposed for securing that the health products or technologies as sold or supplied shall be of that quality; and”

NEW CLAUSES

THAT, the Bill be amended by inserting the following new clauses immediately after clause 27—

Application
product licence.

27A. (1) A person who intends to import, manufacture or sell a health product or technology shall apply to the Authority for the registration of the health product or health technology, in the prescribed form.

(2) An applicant under subsection (1) shall—

- (a) specify the particulars of the person with appropriate knowledge of all aspects of the health product or health technology who shall be responsible for all communication between the applicant and the Authority in the declaration page of the application form; and
- (b) where the applicant is not a citizen of Kenya or is a company incorporated outside Kenya, appoint a local representative who shall be a citizen of Kenya, a person who is or has permanent residence or a company incorporated in Kenya.

(3) The application made under subsection (1) shall be accompanied by—

- (a) a proposed label for use on the health product or technology;
- (b) a copy of the manufacturing licence of the health product or technology, where applicable;

- (c) a copy of the good manufacturing practice certificate from the Authority and the regulatory authority of the country where the health product or technology is manufactured;
- (d) a copy of a certificate of analysis from a quality control laboratory recognized by the Authority, where applicable;
- (e) a copy of the marketing authorization or certificate of registration of the health product or technology from the regulatory authority of the country where the health product or technology is sold;
- (f) the available data on the quality, safety, efficacy and performance of the health product or technology submitted in a common technical dossier format;
- (g) a sample of the health product or technology;
- (h) proof of ownership of the site for the manufacture of the health product or technology, where applicable;
- (i) where the applicant is not a citizen of Kenya or is a company incorporated outside Kenya, a copy of the agreement appointing the local representative;
- (j) where the application relates to a health product or technology which is registered with a foreign regulatory body—
 - (i) a copy of the certificate of registration;
 - (ii) the professional information relating to the health product or technology; and
 - (iii) the conditions of the registration of the health product or technology;
- (k) proof that the applicant holds—
 - (i) a valid practising licence issued by the body responsible for the profession of pharmacy;
 - (ii) a valid wholesale dealer's licence issued in accordance with this Act;
 - (iii) a valid licence to sell poisons issued in accordance with this Act; or
 - (iv) a valid manufacturing licence issued in accordance with this Act; and
 - (v) proof of payment of the application fees as prescribed by the Authority.

- (4) An applicant shall notify the Authority of any variation to the agreement appointing the local representative within seven days of the variation.

Processing of
application for
registration of health
product or
technology.

27B. (1) The Authority shall consider the application made under section 27A, and, shall, if it is satisfied of the safety, efficacy, quality, performance and economic value of the health product or technology, register the health product or technology and issue a certificate of registration in the prescribed form.

(2) The Authority may, while considering the application, approve the details as supplied by the applicant or approve it with such amendments as it may consider appropriate in respect of the following particulars—

- (a) the name under which the health product or technology may be sold;
 - (b) the labelling of the health product or technology;
 - (c) the statement of the representations to be made for the promotion of the health product or technology regarding—
 - (i) the claim to be made for the health product or technology;
 - (ii) the route of administering the health product or technology;
 - (iii) the dosage of the health product or technology;
 - (iv) the storage conditions of the health product or technology;
 - (v) the contra-indications, the side effects and precautions, if any of the health product or technology; and
 - (vi) the package size of the health product or technology.
- (3) When evaluating an application, the Authority may—
- (a) subject a sample of the health product or technology to an evaluation by an analyst; and
 - (b) consider the evaluation report of the analyst that has evaluated the health product or technology.
- (4) Where the Authority is not satisfied as to the quality, safety efficacy, performance or economic value of the health product or technology, it may, after providing an opportunity to the applicant to be heard, reject the application and inform the applicant the reasons for rejection in writing.

Registration during
emergency.

27C. (1) The Authority may, where it considers it necessary to protect public health or in the event of a threat to life or health, issue a provisional certificate of registration for a health product or technology.

(2) A person who intends to obtain the provisional certificate of registration for a health product or technology under subsection (1) shall apply to the Authority in the prescribed form.

(3) Where the applicant is not a citizen of Kenya or is a company incorporated outside Kenya, the applicant shall appoint a local representative who shall be a citizen of Kenya, a person who is or has permanent residence or a company incorporated in Kenya.

(4) An application under subsection (2) shall be accompanied by—

- (a) such documents as may be necessary to support the application;
- (b) where the applicant is not a citizen of Kenya or is a company incorporated outside Kenya, a copy of the agreement appointing the local representative;
- (c) proof that the applicant holds—
 - (i) a valid practising licence issued by the body responsible for the profession of pharmacy;
 - (ii) a valid wholesale dealer's licence issued in accordance with this Act;
 - (iii) a valid licence to sell health products or technologies issued in accordance with this Act;or
 - (iv) a valid manufacturing licence issued in accordance with this Act; and
 - (v) proof of payment of the application fees as prescribed by the Authority.

(5) When determining an application under this section, the Authority shall consider the facts established from the valid marketing authorization for the health product or technology and the report on the assessment of the health product or technology obtained from the authority competent for health products and technologies, if available.

(6) The person to whom the certificate of registration is issued under this section, shall be responsible for the labelling, packaging, advertising and pharmacovigilance system of the health product or technology.

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(7) A provisional certificate of registration issued under subsection (1) shall be valid for two years from the date of issue or until the declaration made under section 35 of the Public Health Act is revoked.

(8) Any variation to the agreement appointing the local representative to the application made under subsection (2) shall be notified to the Authority within seven days of the variation.

Authorization of
unregistered health
product or
technology.

27D. (1) The Authority may, in writing, authorize a person to import or distribute for a specified period to a specified person or institution a specified quantity of a particular health product or technology that is not registered.

(2) A health product or technology distributed pursuant to authorization granted under subsection (1) may be used for such purposes and in such manner and during such period as the Authority may in writing determine.

(3) A person who intends to obtain the authorization under subsection (1), for purposes other than a clinical trial, shall apply to the Authority in the prescribed form.

(4) Where the applicant is not a citizen of Kenya or is a company incorporated outside Kenya, the applicant shall appoint a local representative who shall be a citizen of Kenya, a person who is or has permanent residence or a company incorporated in Kenya.

(5) The application made under subsection (3) shall be accompanied by—

- (a) a product brochure containing relevant chemical, pharmaceutical, pre-clinical pharmacological and toxicological data and where applicable, human pharmacological and clinical data related to the health product or technology for which authority is sought;
- (b) written consent of the applicant, where applicable;

- (c) details of registration or pending registration of the health product or technology with any other regulatory authority, where applicable;
- (d) evidence of compliance by the manufacturer of the health product or technology with good manufacturing practice standards as determined by the Authority;
- (e) reasons why a registered health product or technology cannot be used;
- (f) where the applicant is not a citizen of Kenya or is a company incorporated outside Kenya, a copy of the agreement appointing the local representative;
- (g) proof that the applicant holds—
 - (i) a valid practising licence issued by the body responsible for the profession of pharmacy;
 - (ii) a valid wholesale dealer's licence issued in accordance with this Act;
 - (iii) a valid licence to sell health products or technologies issued in accordance with this Act;
 - or
 - (iv) a valid manufacturing licence issued in accordance with this Act; and
 - (v) proof of payment of the application fees as prescribed by the Authority.

(6) Where the Authority issues an authorization under subsection (1), the person to whom the authorization is issued shall submit to the Authority—

- (a) progress reports after every six months from the date of issuance of the authorization;
- (b) any adverse event report, where an adverse event occurred; and
- (c) a progress report within thirty days after the completion or termination of the use of the health product or technology.

(7) The Authority may, where it is of the opinion that the safety of any patient is compromised or where the scientific reasons for administering the unregistered health product or technology have changed—

- (a) impose any additional conditions;
- (b) request additional information;

- (c) inspect the site where the unregistered health product or technology is manufactured, stored or administered; or
- (d) withdraw the authorization to treat the patient.

(8) The Authority may, by notice in writing withdraw the authorization issued under subsection (1) if any of the purposes or the manner specified in subsection (2) is contravened.

(9) A health product or technology authorized under this section shall be labelled in accordance with this Act.

(10) An applicant shall notify the Authority of any variation to the agreement appointing the local representative within seven days of the variation.

(11) The requirements in this section shall apply to applications for donations of health products and technologies.

CLAUSE 28

THAT, Clause 28 of the Bill be amended—

- (a) in the marginal note by deleting the words “medicines register” and substituting therefor the words “health products and technologies register”;
- (b) in sub-clause (1) by deleting the words “medicines register” and substituting therefor the words “health products and technologies register”; and
- (c) in sub-clause (2) by deleting the words “medicines register” and substituting therefor the words “health products and technologies register”.

CLAUSE 29

THAT, Clause 29 of the Bill be amended—

- (a) in the marginal note by deleting the words “medicines and medical devices” and substituting therefor the words “health products and technologies”;
- (b) by deleting sub-clause (1) and substituting therefor the following new sub-clause (1)—

“(1) Every application for registration of a health product or technology shall be submitted to the Registrar in the prescribed form and shall be accompanied by the prescribed particulars and samples of the relevant health product or technology and by the prescribed registration fee.”

- (c) in sub-section (2) by deleting the phrase “Essential Medicines List or Essential Veterinary Medicines List” and substituting therefor the phrase “Kenya Essential Medicines List, Kenya Essential Diagnostics List, Kenya Essential Medical Supplies List and Kenya Essential Veterinary Medicine List”;
- (d) in sub-clause (3) by deleting the word “medicine” wherever it appears and substituting therefor the words “health product or technology”;
- (e) in sub-clause (4) by—
 - (i) deleting the word “medicine” appearing in the opening sentence and substituting therefor the words “health product or technology”;
 - (ii) deleting paragraph (b) and inserting the following new paragraph—

“(b) that the applicant may, within a period of three months after the date of the notification, furnish the Registrar with the comments on the Authority’s reasons for not being so satisfied.”
- (f) by deleting sub-clause (6) and substituting therefor the following new sub-clause (6)—

“(6) Where the Authority has approved the registration of any health product or technology if it is satisfied of the safety, efficacy, quality, performance and economic value of the health product or technology, the Registrar shall register that health product or technology and shall enter in the register such particulars in regard to the health product or technology as are required by this Act to be so entered and shall issue to the applicant a certificate of registration in the prescribed form in respect of that health product or technology.”
- (g) in sub-clause (7) by deleting the word “medicine” and substituting therefor the words “health product or technology”;
- (h) in sub-clause (8) by deleting the word “medicine” wherever it appears and substituting therefor the words “health product or technology”;
- (i) in sub-clause (9) by deleting the word “medicines” and substituting therefor the words “health products and technologies”;
- (j) in sub-clause (10) by deleting the word “medicine” and substituting therefor the words “health product or technology”;
- (k) in sub-clause (11) by deleting the word “medicine” and substituting therefor the words “health product or technology”;

- (l) in sub-clause (12) by deleting the word “medicine” appearing in the opening sentence and substituting therefor the words “health product or technology”;
- (m) in sub-clause (14) by—
- (i) deleting paragraph (a) and substituting therefor the following new paragraph (a) —
 “(a) Kenya Essential Medicines List, Kenya Essential Diagnostics List and Kenya Essential Medical Supplies List means the list of essential medicines, diagnostics and medical supplies included in the latest editions of the official publications relating to guidelines for standard treatment which is compiled by the state department responsible for Health;”
 - (ii) inserting the word “Kenya” immediately before the phrase “Essential Veterinary Medicines List” appearing in paragraph (b).

NEW CLAUSES 29A & 29B

THAT, the Bill be amended by inserting the following new clauses immediately after clause 29—

Authorization of health products and technologies.

29A. (1) A person shall not import any health product or technology unless—

- (a) the imported health product or technology has been authorized through issuance of an import permit or a written authorization by the Authority; and
- (b) the imported health product or technology is inspected and verified by an inspector of the Authority at the ports of entry prior to its release.

(2) A batch or lot of any registered product shall not be released by the manufacturer prior to the completion of tests for conformity with standards applicable to such product and official batch or lot release by the Authority in cases of biological therapeutics.

(3) Each applicable test conducted by the manufacturer under subsection (2) shall be made on each batch or lot after completion of all processes of manufacture and such test may affect compliance with the standard applicable to the product.

(4) The manufacturer or marketing authorization holder of any registered biological therapeutic shall submit lot summary protocol for each lot that contains registered tests and results of tests

performed and, such manufacturer or marketing authorization holder may be required to submit samples of product from the specified lot to the Authority for official batch or lot release in accordance with the prescribed regulations.

(5) Every batch or lot of a registered biological therapeutic imported into Kenya or manufactured in Kenya shall be evaluated and, on being satisfied of conformity with prescribed standards and payment of prescribed fees, the Director-General shall approve its release into the market and issue a certificate of official batch or lot release in the prescribed format.

(6) The Authority may recognize and accept official lot release certificates issued by other national regulatory authorities of other countries for a specific batch or lots of biological therapeutic manufactured within the territories of those national regulatory authorities, in issuance of a certificate under this section.

(7) A person who contravenes this section commits an offence and shall on conviction be liable—

- (a) in the case of a first offence, to a fine not exceeding one million shillings or to imprisonment for a term not exceeding two years, or to both; or
- (b) in the case of a subsequent offence, to a fine not exceeding two million shillings or to imprisonment for a term not exceeding five years, or to both.

Parallel importation
of health products
and technologies.

No. 17 of 2015.

29B. (1) A person shall not engage in the parallel importation of a health product or technology into Kenya unless—

- (a) the person is incorporated as a limited liability company under the Companies Act;
- (b) the person has been granted a certificate of parallel importation;
- (c) the person is licensed to parallel import the health product or technology;
- (d) the health product or technology has a valid registration in Kenya under this Act; and
- (e) the health product or technology has a valid market authorization in the country of origin.

(2) A person who wishes to undertake parallel importation of a health product or technology shall apply to the Board for a certificate of parallel importation in the prescribed manner.

(3) The Board shall establish and maintain a system that ensures that a registered parallel imported health product or technology can be traced from its sourcing, manufacturing, packaging, storage, transport to its delivery to the health facility, institution or private practice where the health product or technology is intended to be used.

(4) A person who—

(a) is the holder of a certificate of parallel importation or licensee and fails to comply with any requirement or obligation in this Act;

(b) contravenes any prohibition prescribed by the Authority; or

(c) fails to comply with any requirement imposed on that person by the Board pursuant to this Act,

commits an offence and is liable, upon conviction, to a fine not exceeding one million shillings or to imprisonment for a term not exceeding two years, or to both.

CLAUSE 30

THAT, Clause 30 of the Bill be amended—

(a) in sub-clause (1) by deleting the word “medicine” wherever it appears and substituting therefor the words “health product or technology”;

(b) by inserting the following new sub-clause immediately after subclause (2)—

“(2A) A person who makes an application under this section shall provide reasons for the proposed amendments to the register.”

(c) in sub-clause (3), by deleting the word “medicine” wherever it appears in paragraph (b) and substituting therefor the words “health product or technology”.

CLAUSE 31

THAT, Clause 31 of the Bill be amended—

(a) in sub-clause (1) by deleting the word “medicine” and substituting therefor the words “health product or technology”; and

- (b) in sub-clause (3), by deleting the word “medicine” appearing in paragraph (c) and substituting therefor the words “health product or technology”.

CLAUSE 32

THAT, Clause 32 of the Bill be amended—

- (a) by deleting sub-clause (1) and substituting therefor the following new sub-clause (1)—
- “(1) The Authority shall cancel the registration of a health product or technology if—
- (a) a licensee has failed to comply with a condition subject to which a particular health product or technology has been registered;
 - (b) a particular health product or technology does not comply with a prescribed requirement; or
 - (c) it is not in the public interest to make a particular health product or technology available to the public.”
- (b) in sub-clause (2) by deleting the phrase “medicine or medical device” wherever it appears and substituting therefor the phrase “health product or technology”;
- (c) in sub-clause (4)—
- (i) by deleting the words “medicine or medical device” appearing in the opening sentence and substituting therefor the words “health product or technology”; and
 - (ii) by deleting the words “medicine or medical device” appearing in paragraph (b) and substituting therefor the words “health product or technology”; and
- (d) by deleting the words “medicine or medical device” wherever it appears in sub-clause (5) and substituting therefor the words “health product or technology”.

CLAUSE 33

THAT, Clause 33 of the Bill be amended in sub-clause (1) by deleting the words “medicine or medical device” and substituting therefor the words “health product or technology”.

CLAUSE 34

THAT, Clause 34 of the Bill be amended—

- (a) by deleting the words “medicines” and “medicine” wherever they appear and substituting therefor the words “health product or technology”; and
- (b) in the marginal note by deleting the words “medicines” and substituting therefor the words “health products and technologies”.

CLAUSE 35

THAT, Clause 35 of the Bill be amended—

- (a) by deleting the word “medicine” wherever it appears and substituting therefor the words “health product or technology”;
- (b) by deleting sub-clause (1) and substituting therefor the following new sub-clause (1)—
 - “(1) A pharmacist or an enrolled pharmaceutical technologist may, in consultation with the person prescribing the health product or technology and the patient, dispense an interchangeable multi-source health product or technology instead of the health product or technology prescribed by a medical or dental practitioner, nurse or other person registered under the relevant statutes regulating health professionals.”
- (c) in sub-clause (2) by inserting the words “or an enrolled pharmaceutical technologist” immediately after the word “pharmacist”;
- (d) in sub-clause (3) by inserting the words “or an enrolled pharmaceutical technologist” immediately after the word “pharmacist”; and
- (e) in sub-clause (4) by inserting the word “or an enrolled pharmaceutical technologist” immediately after the word “pharmacist”.

CLAUSE 36

THAT, Clause 36 of the Bill be amended—

- (a) in sub-clause (1) by inserting the words “or products” immediately after the words “herbal medicine”; and
- (b) in sub-clause (3) by inserting the phrase “and shall, on conviction be liable to a fine not exceeding one million shillings or imprisonment for a term not exceeding one year, or to both” immediately after the phrase “commits an offence”.

NEW CLAUSE 36A

THAT, the Bill be amended by inserting the following new clause immediately after clause 36—

- Clinical trials.
- 36A.** (1) A health product or technology shall not be used for clinical trial unless an approval is granted by the Authority.
 - (2) An approval under subsection (1) shall only be granted by the Authority after approval by the relevant ethics body has been granted.

(3) A person who intends to commence a clinical trial on a health product or technology shall make an application to the Authority in the prescribed form and the application shall be accompanied by the study protocol in the prescribed format and the prescribed fee.

(4) The study protocol submitted under subsection (2) shall include a post-trial access programme to ensure access of investigational medicinal substances by participants in the trial before grant of marketing authorization by the Authority.

(5) The Authority shall prescribe guidelines for evaluation of applications made under subsection (2) to be implemented for accelerated evaluations during emergency situations, epidemics and outbreaks.

(6) A person granted an approval under this section shall put in place a robust quality assurance system to ensure that the clinical trial is carried out in a manner that ensures the integrity of data generated and the safety and well-being of the participants of the study.

(7) The Authority shall carry out inspection of the clinical trials and monitor compliance of the clinical trials with the prescribed requirements.

(8) Any amendments to clinical trials protocols shall be submitted to the Authority for approval before implementation.

PART V

THAT, the Bill be amended in the title to Part V by deleting the expression “PART V” and substituting therefor the expression “PART IV”.

CLAUSE 37

THAT, Clause 37 of the Bill be amended—

(a) in sub-clause (2) by deleting the words “and dealers in mining, agricultural or horticultural accessories” appearing in paragraph (a);

(b) by inserting the following new sub-clause (3) immediately after sub-clause (2)—

“(3) The Cabinet Secretary shall publish in the *Gazette* the list of scheduled substances prepared under subsection (1).”

(c) by renumbering sub-clause (3) as sub-clause (4);

(d) by deleting sub-clause (4) and substituting therefor the following new sub-clauses

—
“(5) The Authority shall at least once every two years, review the lists under subsection (3), or whenever necessary in the interest of public health and safety.

(6) Any modification of the list of scheduled substances prepared under this section shall be subject to the procedure provided in subsections (1), (2) and (3).”

CLAUSE 38

THAT, Clause 38 of the Bill be amended—

(a) in sub-clause (1) by—

(i) deleting the phrase “the Limitations prescribed by this sub-section” and substituting therefor the phrase “the following limitations”;

(ii) deleting paragraph (c)

(b) by deleting sub-clause (2) and substituting therefor the following new sub-clause (2)—

“(2) A person who is in possession of a scheduled substance otherwise than in accordance with the provisions of this section commits an offence and shall on conviction, be liable to a fine not exceeding two million shillings or to imprisonment for a term not exceeding three years; or to both.”

CLAUSE 39

THAT, Clause 39 of the Bill be amended—

(a) in sub-clause (4) by inserting the word “and” immediately after the words “distribution of the Scheduled Substances”;

(b) by deleting sub-clause (5) and substituting therefor the following new sub-clause (5)—

“(5) A licence issued under this section shall be valid for a period of one year, renewable annually.”

CLAUSE 40

THAT, the Bill be amended by deleting clause 40.

CLAUSE 41

THAT, Clause 41 of the Bill be amended—

- (a) in sub-clause (1)—
 - (i) by deleting paragraph (c);
 - (ii) by deleting paragraph (e);

- (b) in sub-clause (2) —
 - (i) by deleting paragraph (b)
 - (ii) by deleting paragraph (c); and

- (c) by deleting sub-clause (3).

CLAUSE 42

THAT, Clause 42 of the Bill be amended—

- (a) in sub-clause (1) by deleting the expression “paragraph (b) of Section 53(2)” appearing in paragraph (a) and substituting therefor the expression “section 41(2)(b)”;
- (b) in sub-clause (3) by deleting the words “three years” and substituting therefor the words “one year”.

CLAUSE 43

THAT, Clause 43 of the Bill be amended in sub-clause (1)—

- (a) by deleting the opening sentence and substituting therefor the following new opening sentence—

“(1) A qualified healthcare professional may supply or dispense a Scheduled Substance with therapeutic value for the purpose of medical, dental or veterinary treatment, as the case may be, subject to the following provisions—

- (b) in paragraph (b) by—
 - (i) inserting the word “and” immediately after the word “supplied” appearing in sub-paragraph (iii); and
 - (ii) deleting the word “and” appearing in sub-paragraph (iv);

- (c) by deleting paragraph (c).

CLAUSE 44

THAT, Clause 44 of the Bill be amended in sub-clause (3) by deleting the words “two hundred thousand” and substituting therefor the words “five hundred thousand”.

CLAUSE 45

THAT, the Bill be amended by deleting Clause 45 and substituting therefor the following new clause 45—

Automatic
machines.

45. (1) An authorized seller may use an automatic machine to dispense over-the-counter scheduled substances.

(2) The Authority shall develop regulations on the—

- (a) classes of substances permitted;
- (b) quantities of substances to be dispensed;
- (c) records of substances dispensed;
- (d) location of automatic machines; and
- (e) registration of automatic machines.

CLAUSE 46

THAT, the Bill be amended by deleting Clause 46 and substituting therefor the following new clause 46—

Electronic sale of
health products and
technologies.

46. (1) The Authority shall prescribe regulations to provide for the electronic supply and dispensing of scheduled substances including through e-pharmacy, telemedicine, medication therapy management and online pharmacy.

(2) The regulations made under subsection (1) shall provide for—

- (a) licensure of e-pharmacies;
- (b) safety of patients;
- (c) verification of the identity and traceability of patients;
- (d) verification of the identity and traceability of prescribers; and
- (e) integrity, legitimacy and authenticity of prescriptions including avoidance of multiple use of the same prescription.

(3) The electronic supply and dispensing of scheduled substances shall be permitted provided that the supply of such health products and technologies conforms with all requirements for the particular health product or technology in terms of its scheduling status and any other requirements as may be specified in regulations in relation to such supply or dispensing.

(4) In the case of a prescription-only medicine, the required prescription shall have been obtained as a result of at least one physical interaction between an authorised practitioner and the patient within a period of at least six months.

(5) A person who contravenes this section shall be guilty of an offence, and shall on conviction, be liable to a fine not exceeding one million shillings, or to imprisonment for a term not exceeding one year, or to both.

NEW CLAUSE 46A

THAT, the Bill be amended by inserting the following new clause immediately after clause 46—

Dietary
supplements.

46A. (1) A dietary supplement shall—

- (a) have a stated or implied therapeutic purpose; and
- (b) not contain a scheduled substance.

(2) Where a supplement contains a dietary ingredient, the maximum daily dose for the dietary ingredient shall be as per the guidelines prescribed by the Authority.

PART VI

THAT, the Bill be amended in the title of Part VI by deleting the expression “PART VI—MANUFACTURE OF MEDICINAL SUBSTANCES” and substituting therefor the expression “PART V—MANUFACTURE OF HEALTH PRODUCTS”.

CLAUSE 47

THAT, Clause 47 of the Bill be amended—

- (a) in sub-clause (1) by deleting the words “medicinal substance” and substituting therefor the words “health product”;
- (b) by deleting sub-clause (2) and substituting therefor the following new sub-clause (2)—

“(2) A manufacturing licence issued under this section shall be valid for a period of one year, renewable annually.”
- (c) in sub-clause (3) by deleting the words “medicinal substance” and substituting therefor the words “health product”;
- (d) in sub-clause (4) by deleting the words “medicinal substance” and substituting therefor the words “health product”;
- (e) by inserting the following sub-clauses immediately after sub-clause (5)—

“(6) The Authority shall prescribe regulations setting out conditions for the qualifications of personnel involved in the production processes of a health product regulated under this Act.

(7) The personnel qualified to conduct lot release of vaccines and batch release of health products shall submit their qualifications to the Authority.

(8) A person who commits an offence under this section shall on conviction, be liable to a fine not exceeding ten million shillings, or to imprisonment for a term not exceeding ten years, or to both.”

CLAUSE 48

THAT, Clause 48 of the Bill be amended by—

- (a) renumbering the existing provision as sub-clause (1); and
- (b) inserting the following new sub-clauses immediately after the renumbered sub-clause (1)—

“(2) The Authority shall have power to enter and inspect manufacturing premises to confirm compliance with prescribed good manufacturing practices and issue a certificate of compliance in the prescribed format upon payment of prescribed fees.

(3) The Cabinet Secretary shall make regulations for the better carrying out of the provisions of this section.

(4) Without prejudice to the generality of subsection (3), the Cabinet Secretary shall make regulations on—

- (a) revocation and suspension of manufacturing licences;
- (b) withdrawal of revocation of manufacturing licences upon request; and
- (c) transfer of manufacturing licences.”

PART VII

THAT, the Bill be amended in the title of Part VII by deleting the expression “PART VII” and substituting therefor the expression “PART VI”.

NEW CLAUSE 50A

THAT, the Bill be amended by inserting the following new clause immediately after clause 50—

Information that is required to be displayed on the pack.

- 50A.** (1) A person dealing in a therapeutic cosmetic shall indicate—
- (a) the common name of the therapeutic cosmetic;
 - (b) the net weight of the therapeutic cosmetic;
 - (c) all the cosmetic ingredients in the order of prominence but not including flavours or fragrances;
 - (d) the name and address of the manufacturer of the therapeutic cosmetic;
 - (e) a warning statement; and
 - (f) a statement that the therapeutic cosmetic is capable of curing or treating any disease or medical condition.
- (2) The Cabinet Secretary shall make regulations for the effective implementation of this section.
- (3) The regulations made under subsection (1) may—
- (a) require manufacturers of cosmetics to register with the Authority; and
 - (b) impose restrictions, requirements or other conditions on manufacturers of cosmetics, if such restrictions, requirements or conditions are necessary to protect public health.

CLAUSE 51

THAT, Clause 51 of the Bill be amended by inserting the phrase “and shall on conviction be liable to a fine not exceeding one million shillings, or to imprisonment for a term not exceeding two years, or to both” immediately after the word “offence”.

CLAUSE 52

THAT, Clause 52 of the Bill be amended by deleting the phrase “have a therapeutic effect or value shall be treated as a medicine” and substituting therefor the phrase “treat, diagnose or prevent disease, or affect the structure or functions of the body shall be treated as a health product or technology”.

CLAUSE 54

THAT, Clause 54 of the Bill be amended by deleting sub-clause (3) and substituting therefor the following new sub-clause (3)—

“(3) A person who manufactures, sells, supplies, imports or exports a therapeutic cosmetic which contains a prohibited ingredient commits an offence and, shall on conviction, be liable to a fine not exceeding one million shillings, or to imprisonment for a term not exceeding two years, or to both.”

PART VIII

THAT, the Bill be amended in the title of Part VIII by deleting the expression “PART VIII” and substituting therefor the expression “PART VII”.

CLAUSE 55

THAT, Clause 55 of the Bill be amended by deleting sub-clause (1) and substituting therefor the following new sub-clause—

“(1) The Registrar shall keep in the prescribed form a register of all medical devices approved by the Authority.”

CLAUSE 56

THAT, Clause 56 of the Bill be amended by deleting sub-clause (1) and substituted therefor the following new sub-clause—

“(1) A person shall not sell any medical device that is—

- (a) not registered by the Authority;
- (b) adulterated;
- (c) substandard, falsified, falsely labelled or counterfeited; or
- (d) which fails to comply in any way with specifications of this Act or any other law.”

CLAUSE 57

THAT, Clause 57 of the Bill be amended by inserting the phrase “and shall, on conviction be liable to a fine not exceeding two million shillings, or imprisonment for a term not exceeding three years, or to both” immediately after the words “commits an offence”.

CLAUSE 58

THAT, Clause 58 of the Bill be amended—

- (a) in sub-clause (2) by inserting the phrase “in accordance with the most recent World Health Organization’s prescribed guidelines on good manufacturing practice” immediately after the word “Authority”;

- (b) by inserting the following new sub-clauses immediately after sub-clause (2)—

“(3) The Authority shall receive from the Kenya Nuclear Regulatory Authority established under the Nuclear Regulatory Act, 2019 documented evidence of radiation required to enable a medical device perform its therapeutic and diagnostic functions and the intended purpose of the device, for issuance of a registration certificate for a medical device.

(4) An importer, distributor or dealer shall establish and implement documented procedures for the maintenance of importation or distribution records and shall maintain an importation or distribution record of each medical device to be submitted to the Authority.”

CLAUSE 59

THAT, Clause 59 of the Bill be amended—

- (a) in sub-clause (1) by inserting the words “unregistered establishments for medical devices and” immediately after the word “under”; and
- (b) by deleting sub-clause (3) and substituting therefor the following new sub-clause—
 - “(3) A person who sells or supplies unapproved medical devices commits an offence and shall, on conviction be liable to a fine not exceeding one million shillings or to imprisonment for a term not exceeding two years, or to both.”

NEW CLAUSE 59A

THAT, the Bill be amended by inserting the following new clause immediately after clause 59—

Registration of medical devices establishment.

59A. (1) An application for registration of a medical devices establishment shall be submitted to the Authority in the prescribed format and shall be accompanied by the prescribed fees.

(2) An importer, distributor or dealer will establish a system of notification of field safety corrective action and shall notify the Authority of such system.

(3) Where the Authority is satisfied that the application under subsection (1) meets the prescribed requirements, the Director-General shall issue a registration certificate for the medical devices establishment in the prescribed format.

(4) A medical devices establishment registration certificate issued under this section shall be valid for a period of one year, renewable annually upon application in accordance with the prescribed conditions.

(5) The registration certificate for manufacturers shall be valid for five years following a successful reinspection.

(6) The Authority may refuse to issue a medical devices establishment registration certificate where—

- (a) an applicant has made a false or misleading statement in the application;
- (b) the Authority has reasonable grounds to believe that issuing the medical devices establishment registration certificate will constitute a risk to the health or safety of patients, users or other persons; or
- (c) an applicant has failed to meet the prescribed conditions for medical devices establishment registration.

(7) Where the Authority does not issue a medical devices establishment registration certificate under subsection (6), the Authority shall—

- (a) notify the applicant in writing of the reasons for refusing the registration of the establishment; and
- (b) cause the applicant to be notified that the applicant may, within a period of three months from the date of notification, furnish the Authority with additional relevant documentation or evidence in support of the application.

(8) After the issuance of a medical devices establishment registration certificate, where there is a change to any of the information submitted at the time of application, the holder of the registration certificate shall submit the new information to the Authority within ten working days of the change.

NEW PART VIII

THAT, the Bill be amended by inserting the following new Part immediately after the new clause 59A—

PART VIII-THE NATIONAL PHARMACOVIGILANCE SYSTEM

Pharmacovigilance.

59B. (1) The Authority shall establish a National Pharmacovigilance Centre which shall set up and manage the national pharmacovigilance and post marketing surveillance system.

(2) The Centre established under subsection (1) shall receive and maintain all relevant information about suspected adverse drug reactions and adverse events to health products or technologies which have been authorized by the Authority.

(3) The Authority shall conduct both passive surveillance and active surveillance of health products and technologies.

(4) The Authority shall carry out pharmacovigilance audits and inspections in order to ensure compliance with good pharmacovigilance practices and the prescribed requirements.

(5) All entities responsible for placing a health product or technology in the market shall establish and maintain a pharmacovigilance system for managing safety information of health products and technologies.

(6) The entities referred to in subsection (5) shall submit safety information to the Authority in the prescribed manner.

(7) The consumers, general public and health care professionals shall report adverse reactions and adverse events to the Authority in the prescribed manner.

PART XI

THAT, the Bill be amended in the title of Part XI by deleting the expression “PART XI” and substituting therefor the expression “PART IX”.

CLAUSE 60

THAT, the Bill be amended by deleting Clause 60 and substituting therefor the following new clause 60—

Establishment of the
National Quality
Control Laboratory.

60. (1) There is established the National Quality Control Laboratory of the Authority which shall be used as a facility for—

(a) the examination and testing of health products and technologies including vaccines and biopharmaceuticals and any material or substance from or with which and the manner in which drugs may be manufactured, processed

- or treated and ensuring the quality control of drugs and medicinal substances;
- (b) performing chemical, biological, bio-chemical, physiological and pharmacological analysis and other pharmaceutical evaluation;
 - (c) testing, on behalf of the Government, of locally manufactured and imported health products and technologies in the Kenyan market prior to marketing authorization, redistribution and post-distribution;
 - (d) providing technical support to local manufacturers and building their capacity in matters pertaining to quality control of regulated products through on site and off site training and laboratory assessments;
 - (e) conducting investigations into the quality and safety status of regulated products developing and administering a data bank on quality assurance of all health products and technologies and generating scientific evidence and reports on the quality and safety status of the registered products;
 - (f) conducting research and training and providing high quality analytics and expert knowledge in the areas of health products and technologies and active pharmaceutical ingredients; and
 - (g) developing and administering a data bank on quality assurance on behalf of the Authority.

(2) The National Quality Control Laboratory shall be the quality control laboratory of health products and technologies for the Authority.

(3) The Board shall appoint a Director of the National Quality Control Laboratory who shall be responsible to the Authority for the day to day management of the National Quality Control Laboratory.

(4) The Director of the National Quality Control Laboratory shall hold office on such terms and conditions of service as may be specified in the instrument of appointment by the Board.

(5) The Director of the National Quality Control Laboratory shall be a registered pharmacist and shall possess a Master's degree in a science related field from a recognized university.

(6) The Director of the National Quality Control Laboratory shall—

- (a) oversee and coordinate all operations and administration of the National Quality Control Laboratory and provide technical guidance on quality control;
- (b) ensure timely quality control testing of all samples in conformity with national and international standards;
- (c) co-ordinate and supervise the activities of the National Quality Control Laboratory including staff;
- (d) collaborate with other laboratories, regulatory and law enforcement agencies to ensure quality in health products and technologies;
- (e) handle appeals on test results;
- (f) where the laboratory lacks capacity, subcontract laboratory testing services;
- (g) advice the Authority on matters of testing and quality control over health products and technologies; and
- (h) perform any other duties assigned by the Authority from time to time.

(7) The funds to be used for the management of the National Quality Control Laboratory shall consist of all moneys received or recovered under this Part and a portion of the moneys appropriated by the National Assembly to the Authority.

(8) Subject to subsection (7), the monies generated by the National Quality Control Laboratory in the course of the performance of its functions under this section shall be solely expended on the Laboratory.

CLAUSE 61

THAT, Clause 61 of the Bill be amended in sub-clause (1) by deleting the words “Director-General” and substituting therefor the words “Director of the National Quality Control Laboratory”.

PART XII

THAT, the Bill be amended in the title of Part XII by deleting the expression “PART XII” and substituting therefor the expression “PART X”.

CLAUSE 63

THAT, Clause 63 of the Bill be amended—

- (a) in sub-clause (1) by deleting the phrase “medicine, drug, appliance or article” wherever it appears and substituting therefor the words “health product or technology”; and
- (b) in sub-clause (2) by inserting the words “or enrolled pharmaceutical technologists” immediately after the word “pharmacists” appearing in paragraph (d).

CLAUSE 64

THAT, Clause 64 of the Bill be amended by—

- (a) deleting the phrase “a medicine, drug, appliance or article” and substituting therefor the phrase “health product or technology”; and
- (b) deleting the phrase “drug, appliance or article” and substituting therefor the phrase “health product or technology”.

CLAUSE 65

THAT, Clause 65 of the Bill be amended—

- (a) in paragraph (a) by—
 - (i) deleting the words “ or similar article”; and
 - (ii) deleting the word “extravagant,”.
- (b) in paragraph (b) by deleting the word “ an article” and substituting therefor the words “a health product or technology”.

CLAUSE 66

THAT, Clause 66 of the Bill be amended—

- (a) in sub-clause (1) by—
 - (i) deleting the phrase “drug, appliance or article” wherever they appear in paragraph (a) and substituting therefor the phrase “health product or technology”; and
 - (ii) deleting the phrase “medicine, drug, appliance or article” appearing in paragraph (b) and substituting therefor the phrase “health product or technology”;
- (b) in sub-clause (3) by—
 - (i) renumbering the provision as sub-clause (2); and
 - (ii) by inserting the phrase “, enrolled pharmaceutical technologists” immediately after the word “pharmacists” appearing in paragraph (ii).

CLAUSE 67

THAT, Clause 67 of the Bill be amended—

- (a) by deleting the word “articles” appearing in the marginal note and substituting therefor the words “health products and technologies”;
- (b) by deleting sub-clause (1) and substituting the following new sub-clauses—

“(1) Subject to this Act, a person shall not sell by retail a health product or technology consisting of or comprising a substance recommended as a medicine unless there is written so as to be clearly legible on the health product or technology or on a label affixed thereto, or if the health product or technology is sold or supplied in more than one container, on the inner container or on a label affixed thereto—

- (a) the appropriate designation of the substance so recommended or of each of the active constituents, or of each of the ingredients from which it has been compounded; and
- (b) in a case where the appropriate designation of each of the active constituents or ingredients is written, the appropriate quantitative particulars of the constituents or ingredients:

(1A) Subsection (1) shall not apply to a health product or technology made up and supplied for the use of a particular person, being an article prescribed by reference to the needs of that person.”

- (c) in sub-clause (2) by deleting the word “article” wherever it appears in the definition of “appropriate quantitative particulars” and substituting therefor the words “health product or technology”;
- (d) in sub-clause (3) by—
 - (i) deleting the word “an article” appearing in the opening statement and substituting therefor the words “a health product or technology”;
 - (ii) deleting the words “two hundred thousand” appearing in paragraph (a) and substituting therefor the words “one million”;
 - (iii) deleting the words “three hundred thousand” appearing in paragraph (b) and substituting therefor the words “two million”.

CLAUSE 68

THAT, the Bill be amended by deleting Clause 68.

CLAUSE 69

THAT, Clause 69 of the Bill be amended by—

- (a) deleting the word “article” and substituting therefor the words “health product or technology”; and
- (b) deleting the word “articles” and substituting therefor the words “health products and technologies”.

PART XIII

THAT, the Bill be amended in the title to Part XIII by deleting the expression “PART XIII” and substituting therefor the expression “PART XI”.

CLAUSE 71

THAT, Clause 71 of the Bill be amended—

- (a) in the marginal note by deleting the phrase “medicines or medical devices” and substituting therefor the phrase “health products and technologies”; and
- (b) in sub-clause (1) by deleting the phrase “or homoeopathic medicine, preparation or medical device” and substituting therefor the phrase “health products and technologies”.

CLAUSE 72

THAT, Clause 72 of the Bill be amended—

- (a) in the marginal note by deleting the words “medicine or medical devices” and substituting therefor the words “health products and technologies”;
- (b) in sub-clause (1) by—
 - (i) deleting the words “a person” and substituting therefor the words “a registered pharmacist”; and
 - (ii) inserting the phrase “including a health product and technology for emergency use” immediately after the word “technology”; and
- (c) in sub-clause (3) by deleting the words “medicine or medical device product” and substituting therefor the words “health product or technology”.

CLAUSE 73

THAT, Clause 73 of the Bill be amended—

- (a) in the marginal note by deleting the word “goods” and substituting therefor the words “health products and technologies”.
- (b) in sub-clause (1) by deleting the words “drug, article” wherever they appear and substituting therefor the words “health product or technology”;
- (c) in sub-clause (2) by deleting the words “drug or article” wherever they appear and substituting therefor the words “health product or technology”;
- (d) in sub-clause (3) by deleting the words “drug or article” and substituting therefor the words “health product or technology”; and
- (e) in sub-clause (4) by deleting the words “drug or article” and substituting therefor the words “health product or technology”.

CLAUSE 79

THAT, the Bill be amended by deleting Clause 79 and substituting therefor the following new clause 79—

Inspection and
verification of health
products and
technologies at the
ports of entry.

79. (1) A person who imports a health product or technology shall notify the inspectors of the Authority at the ports of entry to conduct pre-clearance inspection and verification.

(2) A person who imports a health product or technology and causes it to be released to the market without inspection and verification under subsection (1) commits an offence.

(3) A person who commits an offence under this section shall on conviction, be liable to a fine not exceeding one million shillings, or to imprisonment for a term not exceeding two years, or to both.

CLAUSE 80

THAT, Clause 80 of the Bill be amended—

(a) in sub-clause (1) by—

(i) deleting the word “article” wherever it appears and substituting therefor the words “health product or technology”; and

(ii) inserting the words “or any other vessel” immediately after the word “vehicle” appearing in paragraph (b).

(b) in sub-clause (6) by deleting the word “article” and substituting therefor the words “health product or technology”;

(c) in sub-clause (7) by deleting the word “article” wherever it appears and substituting therefor the words “health product or technology”;

(d) in sub-clause (8) by deleting the word “article” wherever it appears and substituting therefor the words “health product or technology”;

(e) in sub-clause (9) by deleting the word “article” wherever it appears and substituting therefor the words “health product or technology”;

(f) in sub-clause (10) by deleting the word “article” wherever it appears and substituting therefor the words “health product or technology”;

(g) in sub-clause (11) by deleting the word “article” and substituting therefor the words “health product or technology”;

(h) in sub-clause (12) by deleting the word “article” and substituting therefor the words “health product or technology”.

CLAUSE 81

THAT, the Bill be amended by deleting Clause 81.

CLAUSE 82

THAT, the Bill be amended by deleting Clause 82.

CLAUSE 83

THAT, the Bill be amended by deleting Clause 83.

CLAUSE 85

THAT, Clause 85 of the Bill be amended by deleting the word “article” wherever it appears and and substituting therefor the words “health product or technology”.

CLAUSE 86

THAT, Clause 86 of the Bill be amended in sub-clause (1) by deleting paragraph (b) and substituting therefor the following new paragraph (b)—

“(b) in the case of a subsequent offence, to a fine not exceeding one million shillings, or to imprisonment for a term not exceeding two years, or to both.”

CLAUSE 87

THAT, Clause 87 of the Bill be amended in sub-clause (1) by deleting the word “article” wherever it appears in paragraph (c) and substituting therefor the words “health product or technology”.

PART XIV

THAT, the Bill be amended in the title of Part XIV by deleting the expression “PART XIV” and substituting therefor the expression “PART XII”.

CLAUSE 88

THAT, Clause 88 of the Bill be amended by deleting paragraph (a) and substituting therefor the following new paragraph (a)—

“(a) such monies as may be appropriated by the National Assembly for the purposes of the Authority”.

CLAUSE 90

THAT, Clause 90 of the Bill be amended in sub-clause (2) by deleting the words “think fit” appearing in paragraph (f) and substituting therefor the words “consider appropriate”.

CLAUSE 91

THAT, Clause 91 of the Bill be amended—

- (a) in sub clause (3) by deleting the words “Kenya National Audit Office” and substituting therefor the words “Auditor-General”; and
- (b) in sub clause (4) by deleting the words “Kenya National Audit Office” and substituting therefor the words “Auditor-General”.

CLAUSE 92

THAT, Clause 92 of the Bill be amended in sub-clause (2) by inserting the phrase “, with the approval of the Cabinet Secretary of the National Treasury” immediately after the word “may”.

PART XV

THAT, the Bill be amended in the title of Part XV by deleting the expression “PART XV” and substituting therefor the expression “PART XIII”.

CLAUSE 95

THAT, Clause 95 of the Bill be amended—

(a) in sub-clause 2 by—

- (i) deleting the word “drugs,” in paragraph (a)(i);
- (ii) deleting the words “any drug” in paragraph (a)(ii);
- (iii) deleting the word “product” and substituting therefor the word “products” in paragraph (d);
- (iv) deleting the word “drugs” wherever it appears and substituting therefor the words “health products or technologies” in paragraph (h);
- (v) deleting the word “article” and substituting therefor the words “health product or technology” in paragraph (k);
- (vi) deleting the word “articles” and substituting therefor the words “health products and technologies” in paragraph (m);
- (vii) deleting the words “drugs, medical devices” and substituting therefor the words “health products and technologies” in paragraph (o);
- (viii) deleting the word “medicines” and substituting therefor the words “health products and technologies” in paragraph (v);
- (ix) deleting paragraph (x) and substituting therefor the following new paragraph (x)—

“(x) governing administration of clinical trials of health products and technologies;”

- (x) deleting the words “medicine, medical device” and substituting therefor the words “health product or technology” in paragraph (aa);

- (xi) deleting paragraph (bb) and substituting therefor the following new paragraph— “(bb) providing for the manner in which a pharmacist, an enrolled pharmaceutical technologist or a person otherwise authorized under this Act may dispense health products or technologies”;
- (xii) deleting paragraph (dd) and substituting therefor the following new paragraph (dd)—

“(dd) the compounding of health products and technologies and the dispensing of health products and technologies”

- (xiii) deleting the words “generally, for giving effect to this Act” appearing in paragraph (ii);
- (xiv) inserting the following new paragraphs immediately after paragraph (ii)—
 - “(jj) on pharmacovigilance and post market surveillance;
 - (kk) official regulatory lot release of vaccines and other biological products imported and manufactured in Kenya;
 - (ll) pricing of health products and technologies;
 - (mm) good practices in the regulation of health products and technologies;
 - (nn) inspections, licensure and certification of the manufacture of health products and technologies by health facilities;
 - (oo) inspections, licensure and certification of manufacture of health products and technologies and other regulated products by facilities not directly regulated by the Authority including steel industries, sugar industries;
 - (pp) inspection and recognition of pharmaceutical quality control laboratories;
 - (qq) to regulate licit use of narcotic and psychotropic substances; and
 - (rr) to regulate parallel importation of health products and technologies;”

(b) by renumbering sub-clause (2) as sub-clause (3).

CLAUSE 96

THAT, Clause 96 of the Bill be amended—

(a) in sub-clause (1) by—

- (i) deleting paragraph (d) and substituting therefor the following new paragraph (d)—
 - “(d) all members and staff of the former Board shall be deemed to be members and staff of the Authority, and subject to the provisions of any

rules made under this Act, shall continue in office for the period for which they were appointed as members and staff of the former Board.”

(ii) inserting the following new paragraph immediately after paragraph (d)—

“(e) the staff of the Pharmacy and Poisons Board for the time being working in the directorate responsible for the regulation of health products and technologies shall be deemed to be staff of the Authority, and subject to the provisions of any rules made under this Act, shall continue in office for the period for which they were appointed as staff of the Pharmacy and Poisons Board.”

(b) by deleting the sub-clause (2) and substituting therefor the following new sub-clause (2)—

“(2) In this section, “the former Board” means the Board of the National Quality Control Laboratory established under the Pharmacy and Poisons Act, Cap. 244.”

(c) in sub-clause (3)—

(a) by deleting the word “twelve” appearing in the opening sentence and substituting therefor the words “twenty four”;

(b) by inserting the following new paragraph immediately after paragraph (b)—

“(c) after the expiry of the period of twenty four months—

(i) the Pharmacy and Poisons Board shall be dissolved, and the provisions of subsection (1)(a), (b) and (c) shall, with the necessary modifications, apply; and

(ii) the remaining members and staff of the Pharmacy and Poisons Board shall be deemed to be members and staff of the Authority, and subject to the provisions of any rules made under this Act, shall continue in office for the period for which they were appointed as members and staff of the Pharmacy and Poisons Board.”

CLAUSE 97

THAT, Clause 97 of the Bill be amended by inserting the words “with reference to section 96 (3)” immediately after the words “that Schedule” in sub-clause (1).

SECOND SCHEDULE

THAT, the Bill be amended by deleting the Second Schedule.

THIRD SCHEDULE

THAT, the Bill be amended by deleting the Third Schedule.

FOURTH SCHEDULE

THAT, the Bill be amended by deleting the Fourth Schedule and substituting therefor the following new Schedule—

FOURTH SCHEDULE (s. 21 (2))
SCIENTIFIC ADVISORY COMMITTEES

1. Human Health Products and Technologies Committee.
2. Pharmacovigilance Committee.
3. Cosmetics and Borderline Products Committee.
4. Clinical Trial Scientific Technical Advisory Committee.
5. Dietary Supplements Committee.
6. Digital Health and Technologies Committee.
7. Veterinary Health Products and Technologies Committee.

SEVENTH SCHEDULE

THAT, the Seventh Schedule of the Bill be amended by—

- (a) deleting the word “Board” in the paragraph on Cap. 244
- (b) deleting the phrase “(s. 116) and substituting the phrase (“s.97”).
- (c) deleting the paragraph on Cap. 254.

- 2) **Notice is given that the Member for Mathare (Hon. Anthony Oluoch) intends to move the following amendments to the Kenya Drugs Authority Bill, 2022 at the Committee Stage—**

LONG TITLE

THAT, the Bill be amended by deleting the Long Title and substituting therefor the following new Long Title—

“AN ACT of Parliament to establish the Kenya Health Products and Technologies Authority to ensure safety, quality and efficacy or performance of drugs, poisons, therapeutic and biological products, therapeutic cosmetics, herbal medicines and products, chemical substances, medical devices, veterinary products and other health technologies; to provide for the harmonization and administration of the laws relating to the regulation of, drugs, poisons, therapeutic products, therapeutic cosmetics, chemical products, veterinary products and medical devices and the control and safe handling of poisons; to safeguard the security of the supply chains for, therapeutic products, cosmetics and veterinary products; to provide for measures to optimize the use of therapeutic products in health care in Kenya and for connected purposes.”

CLAUSE 2

THAT, the Clause 2 of the Bill be amended—

(a) in the definition of the term “advertisement” by deleting the words “herbal medicines and products”;

(b) in the definition of the term “article” by—

(i) deleting the words “herbal medicine” appearing in paragraph (a); and

(ii) deleting the words “herbal medicine” appearing in paragraph (b);

(c) in the definition of “authorized seller of scheduled substances” by inserting the words “and enrolled as a pharmaceutical technologist or registered as a pharmacist” immediately after the word “Act”;

(d) in the definition of the term “health products and technologies” by deleting the words “herbal medicines and products”;

(e) by deleting the definition of the term “herbal medicine or product” and substituting therefor the following new definition—

“herbal medicine or product” means a plant derived material or preparations with claimed therapeutic or other health benefits, which contain either raw or processed ingredients from one or more plants or material of inorganic or animal origin and includes herbs, herbal materials, herbal preparations, finished herbal products that contain active ingredients, parts of plants or other plant materials or combinations and excludes herbs, herbal materials, herbal preparations, finished herbal products sold or dispensed on a small scale by traditional health practitioners;”

(f) in the definition of the term “medicine” by inserting the words “other than herbal medicines or products” immediately after the words “or mixture of substances”.

(g) in the definition of “pharmacy” by inserting the words “licensed and” immediately after the words “carried out by” appearing in paragraph (a);

(h) deleting the definition of “chemical substance” and substituting therefor the following new definition—

“chemical substance” means any substance or mixture of substances prepared, sold or represented for use as a germicide, antiseptic, disinfectant, pesticide, insecticide, rodenticide, vermicide, detergent or any other substance or mixture of substances which the Authority may, declare to be a chemical substance;

(i) deleting the definition of “therapeutic cosmetic” and substituting therefor the following new definition—

“therapeutic cosmetic” means a product with the ability to trigger biological actions on the dermis, to target and repair skin issues, to prevent future damage and contains ingredients that are usually not found in regular cosmetics or at higher strengths than could be sold safely over the counter;”.

CLAUSE 3

THAT, Clause 3 of the Bill be amended—

- (a) in sub-clause (1) by deleting paragraph (c); and
- (b) by inserting the following new sub-clause immediately after sub-clause (2)—
 - (3) This Act shall not apply to the regulation of herbal medicines or products.

CLAUSE 6

THAT, Clause 6 of the Bill be amended in sub-clause (4) by deleting the word “ten” in appearing in paragraph (c) and substituting therefor the word “fifteen”.

CLAUSE 8

THAT, Clause 8 of the Bill be amended in sub-clause (7) by inserting the words “, fair representation of persons with disabilities” immediately after the words “regional balance.”

CLAUSE 12

THAT, Clause 12 of the Bill be amended by deleting the words “and herbal drugs” appearing in paragraph (o).

CLAUSE 23

THAT, Clause 23 of the Bill be amended in sub-clause (2) by —

- (a) deleting the words “one million” appearing in paragraph (a) and substituting therefor the words “two million”; and
- (b) deleting the words “two million” appearing in paragraph (b) and substituting therefor the words “five million”.

CLAUSE 29

THAT, Clause 29 of the Bill be amended by deleting sub-clause (9).

CLAUSE 35

THAT, Clause 35 of the Bill be amended in sub-clause (2) by inserting the word “registered” immediately after the words “may prohibit a”.

CLAUSE 36

THAT, the Bill be amended by deleting Clause 36.

CLAUSE 39

THAT, Clause 39 of the Bill be amended in sub-clause (4) by inserting the words “or pharmaceutical technologists” immediately after the words “a Registered Pharmacist”.

CLAUSE 41

THAT, Clause 41 of the Bill be amended in sub-clause (1) by inserting the words “or pharmaceutical technologists” immediately after the words “a pharmacist” appearing in paragraph (b)”.

CLAUSE 42

THAT, Clause 42 of the Bill be amended by—

(a) deleting sub-clause (1) and substituting therefor the following new sub-clause (1)—

“(1) An authorized seller shall enter a record of such particulars of the scheduled substance before delivery of the scheduled substance under this Act.”

(b) inserting the following new sub-clause (2) immediately after the new sub-clause (1)—

“(2) A record under subsection (1) shall be in the format prescribed by the Authority and shall indicate —

- (a) the date of the sale;
- (b) the name and address of the purchaser;
- (c) the quantity of the scheduled substances sold; and
- (d) the purpose for which it is stated by the purchaser to be required.”

(c) renumbering sub-clause (2) as sub-clause (3); and

(d) renumbering sub-clause (3) as sub-clause (4).

CLAUSE 51

THAT, the Bill be amended in clause 51 by inserting the words “and, on conviction, shall be liable to a fine not exceeding one million shillings, or to imprisonment for a term not exceeding two years, or to both” immediately after the word “offence”.

CLAUSE 54

THAT, the Bill be amended by deleting clause 54.

CLAUSE 63

THAT, Clause 63 of the Bill be amended by deleting sub-clause (3).

- 3) **Notice is given that the Member for Suba North (Hon. Millie Odhiambo-Mabona) intends to move the following amendments to the Kenya Drugs Authority Bill, 2022 at the Committee Stage—**

CLAUSE 1

THAT, Clause 1 of the Bill be amended by deleting the phrase “Kenya Drugs Authority Act, 2022” and substituting therefor the phrase “Kenya Drug and Health Technologies Act, 2022”.

CLAUSE 2

THAT, Clause 2 of the Bill be amended—

- (q) in the definition of the word “drug” by inserting the words “herbal medicine” immediately after the words “any medicine” wherever it appears ;
- (r) in the definition of the term “falsified medicines” by inserting the phrase “;” immediately after the words “of active or other ingredients”;
- (s) by deleting the definition of the term “health products and technologies” and substituting therefor the following new definitions—

“health products” means chemical substances, therapeutic cosmetics, herbal medicines, medicines, scheduled substances and related products and substances; and

“health technologies” means medical devices including radiation-emitting devices and related products;

- (t) by deleting the definition of the term “herbal medicine or product” and substituting therefor the following new definitions—

“herbal medicine” means the use of plants to treat disease and enhance general health and wellbeing; and

“herbal product” means a plant derived material or preparations with claimed therapeutic or other human or veterinary health benefits, which contain either raw or processed ingredients from one or more plants, or material of inorganic or animal origin;

- (u) in the definition of the term “manufacture” by deleting the words “making a product or medicinal substance and includes” and substituting therefor the words “making a medicinal substance or product and includes extracting”
- (v) in the definition of the term “registered midwife” by deleting the words “by law to practice the profession of midwife in Kenya” and substituting therefor the words “to practice as such under the Nurses and Midwives’ Act”;
- (w) in the definition of the term “substandard medicines” by inserting the words “under this Act or any other written law” immediately after the words “defined specifications”;

- (x) in the definition of the term “therapeutic cosmetic” by deleting the words “or altering the complexion”
- (y) inserting the following definition in the proper alphabetical sequence—

“alternative medicine” has the meaning assigned to it under the Health Act, 2017;

CLAUSE 7

THAT, Clause 7 of the Bill be amended by —

- (a) deleting paragraph (a) and substituting therefor the following new paragraph (a)—
 - “(a) is a state officer”;
- (b) deleting paragraph (b); and
- (c) deleting paragraph (e) and substituting therefor the following new paragraph (e)—
 - “(e) is a public officer”.

CLAUSE 8

THAT, Clause 8 of the Bill be amended—

- (a) in sub-clause (6) by—
 - (i) deleting paragraph (a) and substituting therefor the following new paragraph (a)—
 - “(a) is a state officer”;
 - (ii) deleting paragraph (b);
- (b) in sub-clause (7) by deleting the words “need for regional balance” and substituting therefor the words “need for youth representation, regional and ethnic balance”.

CLAUSE 10

THAT, Clause 10 of the Bill be amended in sub-clause (1) by deleting the words “permission of the” appearing in paragraph (f) and substituting therefor the words “notifying the”.

CLAUSE 12

THAT, Clause 12 of the Bill be amended by—

- (a) deleting the word “ensure” appearing in paragraph (a) and substituting therefor the word “set”;

(b) deleting paragraph (b) and substituting therefor the following new paragraph (b)—

“(b) ensure that there is compliance with existing legislation through a process of active inspection and investigation”;

(c) deleting the word “being” appearing in paragraph (g); and

(d) inserting the words “chemicals, medicine” immediately after the words “rational use of” appearing in paragraph (o).

CLAUSE 13

THAT, Clause 13 of the Bill be amended by deleting the words “co-opt in such committees” appearing in paragraph (f) and substituting therefor the words “hire as consultants such”.

CLAUSE 17

THAT, Clause 17 of the Bill be amended in sub-clause (2) by inserting the words “youth inclusion” immediately after the words “The principles of”.

CLAUSE 21

THAT, Clause 21 of the Bill be amended by—

(a) inserting the following new sub-clause immediately after sub-clause (9)—

“(9A) The Board shall prepare and submit annually a report of its activities including that of any subcommittees to the Cabinet Secretary and the Cabinet Secretary shall submit the said report to parliament.”

CLAUSE 22

THAT, Clause 22 of the Bill be amended —

- (a) in sub-clause (2) by deleting the words “five years” appearing in paragraph (b) and substituting therefor the words “three years”;
- (b) by deleting sub-clause (3).

CLAUSE 24

THAT, Clause 24 of the Bill be amended —

- (a) in sub-clause (3) by inserting the following new paragraph (c)—
“(c) is a herbal medicine”;

(b) in sub-clause (4) by—

- (i) deleting the words “one hundred” appearing in paragraph (a) and substituting therefor the words “five hundred”; and
- (ii) deleting the words “two hundred thousand shillings or to imprisonment for a term not exceeding five years” appearing in paragraph (b) and substituting therefor the words “one million shillings or to imprisonment for a term not exceeding three years”.

CLAUSE 27

THAT, Clause 27 of the Bill be amended by—

(a) inserting the following new sub-clause (1)—

“(1) The Authority may issue product licences as provided under this Act.”

(b) renumbering the existing sub-clause (1) as sub-clause (2).

CLAUSE 28

THAT, Clause 28 of the Bill be amended in—

- (a) sub-clause (1) by inserting the words “and herbal medicines” immediately after the word “medicines”;
- (b) sub-clause (2) by inserting the words “and herbal medicines” immediately after the word “medicines”.

CLAUSE 29

THAT, Clause 29 of the Bill be amended—

(a) by inserting the following new sub-clause (1)—

“(1) Any pharmacist may apply for the registration of a medicine, herbal medicine or medical device as provided for under this Act.”

- (b) in the existing sub-clause (1) by inserting the words “herbal medicine” immediately after the word “medicine”;
- (c) in sub-clause (3) by inserting the words “or herbal medicine” immediately after the word “medicine” wherever it appears;
- (d) in sub-clause (4) by inserting the words “or herbal medicine” immediately after the word “medicine”;
- (e) in sub-clause (6) by inserting the words “or herbal medicine” immediately after the word “medicine” wherever it appears;
- (f) in sub-clause (7) by inserting the words “or herbal medicine” immediately after the word “medicine”;
- (g) in in sub-clause (8) by inserting the words “and herbal medicine” immediately after the word “medicine” wherever it appears;

(h) in sub-clause (9) by—

(i) inserting the words “and herbal medicine” immediately after the word “medicine”;

(ii) inserting the words “under this Act or any other written law” immediately after the words “already registered”;

(i) in sub-clause (12) by inserting the words “and herbal medicine” immediately after the word “medicine”;

(j) by inserting the following new sub-clauses immediately after sub-clause 12 —

“(12A) The Authority may reject any application if the applicant fails to meet the standards as required by this Act or any other written law.

(12B) A person dissatisfied with the decision of the Registrar may appeal to the Board within sixty days.

(12C) Where a person is dissatisfied with the decision of the Board, the applicant may appeal to the High Court within thirty days from the date of the decision being communicated to him or her.”

(k) in the existing sub-clause (13) by deleting the words “appropriate period referred to in sixty days” and substituting therefor the words “ appropriate period of appeal”.

CLAUSE 30

THAT, Clause 30 of the Bill be amended—

(a) in sub-clause (1) by inserting the words “or herbal medicine” immediately after the word “medicine” wherever it appears;

(b) in sub-clause (3) by inserting the words “or herbal medicine” immediately after the word “medicine” wherever it appears in paragraph (b);

(c) by inserting the following new sub-clause (3)—

“(3) The applicant shall provide reasons for the proposed amendments.”

(d) renumbering the existing sub-clause (3) as sub-clause (4).

CLAUSE 31

THAT, Clause 31 of the Bill be amended in sub-clause (1) by inserting the words “or herbal medicine” immediately after the word “medicine”.

CLAUSE 32

THAT, Clause 32 of the Bill be amended—

- (a) in sub-clause (1), by inserting the words “herbal medicine” immediately after the word “medicine” wherever it appears;
- (b) in sub-clause (2), by inserting the words “herbal medicine” immediately after the word “medicine” wherever it appears;
- (c) in sub-clause (4) by inserting the words “herbal medicine” immediately after the word “medicine” wherever it appears;
- (d) in sub-clause (5) by inserting the words “herbal medicine” immediately after the word “medicine” wherever it appears.

CLAUSE 33

THAT, Clause 33 of the Bill be amended—

- (a) in sub-clause (1) by inserting the words “herbal medicine” immediately after the word “medicine” wherever it appears;
- (b) in sub-clause (2) by inserting the words “herbal medicine” immediately after the word “medicine” wherever it appears;
- (c) by inserting the following new sub-clause immediately after sub-clause (2)—

“(2A) In the case of cancellation of registration of a herbal medicine, the Registrar shall in such case specify-

- (a) the name under which the herbal medicine is registered;
- (b) the active components of the herbal medicine;
- (c) the name of the applicant;
- (d) the name of the person who has propriety rights over the herbal medicine;
- (e) the registration number allocated to the herbal medicine; and
- (f) the conditions if any, subject to which that medicine is registered.

CLAUSE 35

THAT, Clause 35 of the Bill be amended in sub-clause (4) by inserting the following new paragraph (d)—

“(d) unless the purchaser or patient is first informed of the same and agrees to the change”.

CLAUSE 36

THAT Clause 36 of the Bill be amended by deleting the words “on a commercial scale” appearing after the words “A person who” in sub-clause (1).

CLAUSE 40

THAT, Clause 40 of the Bill be amended—

- (a) in sub-clause (4) by deleting the words “whose decision thereon shall be final”:
- (b) in sub-clause (7) by deleting the words “two hundred thousand shillings or to imprisonment for a term not exceeding two years” and substituting therefor the words “one million shillings or to imprisonment for a term not exceeding three years”.

CLAUSE 41

THAT, Clause 41 of the Bill be amended in sub-clause (1) by inserting the words “enrolled pharmaceutical technologist and registered pharmacist” immediately after the word “practitioner” appearing in paragraph (d)

CLAUSE 42

THAT, Clause 42 of the Bill be amended—

- (a) in sub-clause (2) by deleting the words provided that where a person represents that he urgently requires a Schedule Substances for the purpose of his trade, business or profession and satisfies the seller that by reason of some emergency he is unable before delivery to furnish the order in writing, the seller may forthwith deliver the Scheduled Substances to the purchaser who will within twenty four hours of the sale furnish the seller with the written order” appearing in paragraph (a);
- (b) in sub-clause (3) by deleting the words “one hundred thousand” and substituting therefor the words “one million”

CLAUSE 43

THAT, Clause 43 of the Bill be amended in sub-clause (3) by deleting the words “one hundred thousand” and substituting therefor the words “one million”.

CLAUSE 44

THAT, Clause 44 of the Bill be amended in sub-clause (3) by deleting the words “two hundred thousand” and substituting therefor the words “one million”.

CLAUSE 45

THAT, Clause 45 of the Bill be amended by deleting the words “five hundred thousand” and substituting therefor the words “one million”.

CLAUSE 46

THAT, Clause 46 of the Bill be amended by deleting the word “This” and substituting therefor the words “The electronic supply of medicine”.

CLAUSE 60

THAT, Clause 60 of the Bill be amended in sub-clause (1) (c) by inserting the words “medicinal herbs” immediately after the words “medicinal substances” wherever it appears.

CLAUSE 63

THAT, Clause 63 of the Bill be amended in sub-clause (1) by inserting the words “medicinal herbs” immediately after the word “medicine” wherever it appears.

CLAUSE 68

THAT, Clause 68 of the Bill be amended in sub-clause (1) by inserting the words “or herbal medicine” immediately after the word “medicine”.

CLAUSE 71

THAT, Clause 71 of the Bill be amended in sub-clause (1) by inserting the words “or herbal medicine” immediately after the word “medicine”.

CLAUSE 72

THAT, Clause 72 of the Bill be amended in sub-clause (3) by inserting the words “herbal medicine” immediately after the word “medicine”.

CLAUSE 95

THAT, Clause 95 of the Bill be amended in sub-clause 2 by—

- (a) inserting the words “and herbal medicines” immediately after the word “medicines” appearing in paragraph (v);
- (b) inserting the words “herbal medicine” immediately after the word “medicine” appearing in paragraph (bb);
- (c) inserting the words “herbal medicines” immediately after the word “medicines” appearing in paragraph (bb);
- (d) deleting paragraph (dd) and substituting therefor the following new paragraph (dd)—

“(dd) the compounding of medicines and herbal medicines and the dispensing of medicines, herbal medicines and medical devices”.

FOURTH SCHEDULE

THAT, the Fourth Schedule of the Bill be amended—

- (a) in paragraph 1(1) by deleting the words “Cabinet Secretary” and substituting therefor the word “Board”; and
- (b) in paragraph 2(1) by inserting the words “and herbal medicines” immediately after the words “human medicinal products”.

- 4) Notice is given that the Member for Homa Bay Town (Hon. Peter Kaluma) intends to move the following amendments to the Kenya Drugs Authority Bill, 2022 at the Committee Stage—

CLAUSE 2

THAT, Clause 2 of the Bill be amended by—

- (z) deleting the definition of “veterinary medicine”;
- (aa) deleting the definition of “enrolled pharmaceutical technologist” and substituting therefor the following new definition—

“enrolled pharmaceutical technologist” means a pharmaceutical technologist whose name appears on the Roll;”

- (bb) by deleting the words “or material of inorganic or animal origin” in the definition of “herbal medicine or product”.

CLAUSE 3

THAT, Clause 3 of the Bill be amended by deleting sub-clause (2).

CLAUSE 6

THAT, Clause 6 of the Bill be amended in—

- (a) sub-clause (4) by inserting the words “established for regulation of pharmacy, medicine or engineering profession” immediately after the word “body” appearing in paragraph (d);
- (b) deleting sub-clause (6);
- (c) deleting sub-clause (8); and
- (d) deleting sub-clause (9).

CLAUSE 7

THAT, Clause 7 of the Bill be amended —

- (d) by deleting paragraph (a);
- (e) by deleting paragraph (b); and
- (f) in paragraph (d) by deleting the word “is” and substituting therefor the words “has been”.

CLAUSE 8

THAT Clause 8 of the Bill be amended—

- (a) in sub-clause (2) by inserting the words “medical practitioner or medical engineer” immediately after the word “pharmacist” appearing in paragraph (a)(ii);
- (b) in sub-clause (6)—
 - (i) by deleting paragraph (a) and (b); and
 - (ii) in paragraph (e) by deleting the word “is” and substituting therefor the words “has been”;

- (c) in sub-clause (7) by deleting the words “regard shall be had of the need for regional balance and the realisation of the principle that at least one third of the members must be from either gender” and substituting therefor the words “regard shall be had to the need for ethnic and regional balance and the need to ensure that person of same biological sex shall not comprise more than two thirds of the members of the Board”.

CLAUSE 10

THAT, Clause 10 of the Bill be amended by deleting sub-clause (2).

CLAUSE 13

THAT, Clause 13 of the Bill be amended in paragraph (c) by deleting the word “an” and substituting therefor the word “lawful”.

CLAUSE 21

THAT, the Bill be amended by deleting Clause 21

CLAUSE 22

THAT, Clause 22 of the Bill be amended by inserting a new sub-clause (4) as follows—

“(4) Subsection (1) shall not apply to traditional medicines or products.”

CLAUSE 31

THAT, Clause 31 of the Bill be amended in sub-clause (1) by deleting the words “who is duly licensed to practice the profession of pharmacy and holds a valid practising certificate to apply for the registration of a medicine”.

CLAUSE 32

THAT, Clause 32 of the Bill be amended by deleting sub-clause (5).

CLAUSE 38

THAT, Clause 38 of the Bill be amended in sub-clause (1) by—

- (a) deleting the words “on premises registered by the Authority” appearing in paragraph (b);
- (b) deleting the words “for mining, agricultural or horticultural purposes” appearing in paragraph (c); and
- (c) deleting the words “by a qualified medical practitioner, dentist or veterinary surgeon or by a hospital, dispensary or similar institution” appearing in paragraph (e);

CLAUSE 39

THAT, Clause 39 of the Bill be amended in sub-clause (4) by inserting the words “a qualified pharmacist, medical practitioner or medical engineering practitioner or holder of diploma in pharmacy, pharmaceutical technology” immediately after the words “holding the licence is”.

CLAUSE 41

THAT, Clause 41 of the Bill be amended—

(d) in sub-clause (1) by—

- (i) inserting the words “or pharmaceutical technologist or dispensing chemist” immediately after the word “pharmacist” appearing in paragraph (b);
- (ii) deleting the word “or veterinary” appearing in paragraph (d); and
- (iii) by deleting the words “by an order whether general or special, of the Cabinet Secretary: but it shall be an offence to sell Scheduled substances to any of the persons or institutions specified in paragraphs (d) and (f) unless a registered pharmacist is in direct control of the scheduled substances at the premises from which they are sold”

(e) in sub-clause (2) by deleting the words “or veterinary surgeon’ appearing in paragraph (a);

CLAUSE 43

THAT, Clause 43 of the Bill be amended in sub-clause (1) by deleting the words “or veterinary surgeon’.

CLAUSE 46

THAT, the Bill be amended by deleting the words “This” and substituting therefor the words “Electronic sale of medicines”.

CLAUSE 81

THAT, the Bill be amended by deleting the words “or the Director of Veterinary Services, in relation to any matter appearing to affect the general interests of agriculture in Kenya”.

CLAUSE 95

THAT, Clause 95 of the Bill be amended by in sub-clause 2 by deleting the words “or veterinary surgeon’ appearing in paragraph (aa).

FOURTH SCHEDULE

THAT, the Bill be amended by deleting the Fourth Schedule

SEVENTH SCHEDULE

THAT, the Seventh Schedule of the Bill be amended by deleting the paragraph on No. 14 of 1994.

- 5) **Notice is given that the Member for Rarieda (Hon. (Dr.) Otiende Amollo) intends to move the following amendments to the Kenya Drugs Authority Bill, 2022 at the Committee Stage—**

CLAUSE 2

THAT, the Clause 2 of the Bill be amended by deleting the definition of “pharmacy” and substituting therefor the following new definition—

Cap. 244. “pharmacy” has the meaning assigned to it under the Pharmacy and Poisons Act.

CLAUSE 8

THAT, the Clause 8 of the Bill be amended in sub-clause (2) by deleting paragraph (j) and substituting therefor the following new paragraph—

“(j)one person, not being a Governor, who is a registered pharmacist of good standing, nominated by the Council of Governors”;

CLAUSE 39

THAT, the Clause 39 of the Bill be amended in the marginal note by deleting the words “Wholesale Dealer’s Licence” and substituting therefor the word “Local Technical Representative Licence”.



LIMITATION OF DEBATE

The House resolved on Wednesday, February 14, 2024 as follows—

Limitation of Debate on the Annual Estimates and Committee of Supply

- II. THAT**, each speech in a **debate on the Report of the Budget & Appropriations Committee on Budget Estimates** contemplated under Standing Orders 239 and 240 be limited as follows:
- (i) **General Supply Debate**:- A maximum of three (3) sitting days with thirty (30) minutes for the Mover in moving and fifteen (15) minutes in replying; a maximum of fifteen (15) minutes for each of the Chairpersons of the Departmental Committees and a maximum of ten (10) minutes for any other Member speaking, except the Leader of the Majority Party and the Leader of the Minority Party who shall be limited to a maximum of fifteen (15) minutes each; and that priority in speaking be accorded to the Leader of the Majority Party, the Leader of the Minority Party and the respective Chairpersons of the Departmental Committees in the order that they appear in the Second Schedule to the Standing Orders; and,
 - (ii) **Committee of Supply**: - A maximum of six (6) sitting days for the consideration of the proposed allocations to the respective Votes/Programmes in the order specified in the Schedule submitted by the Budget and Appropriations Committee.

Limitation of Debate on Bills sponsored by Parties or Committees

- III. THAT**, each speech in a debate on **Bills sponsored by a Committee, the Leader of the Majority Party or the Leader of the Minority Party** shall be limited as follows: A maximum of forty five (45) minutes for the Mover, in moving and fifteen minutes (15) in replying, a maximum of thirty (30) minutes for the Chairperson of the relevant Committee (if the Bill is not sponsored by the relevant Committee), and a maximum of ten (10) minutes for any other Member speaking, except the Leader of the Majority Party and the Leader of the Minority Party, who shall be limited to a maximum of fifteen minutes (15) each (if the Bill is not sponsored by either of them); and that priority in speaking shall be accorded to the Leader of the Majority Party, the Leader of the Minority Party and the Chairperson of the relevant Departmental Committee, in that order.

Limitation of Debate on Other Committee Reports

- IV. THAT**, each speech in a debate on **Other Committee Reports**, including a Report of a Joint Committee of the Houses of Parliament or any other Report submitted to the House for which limitation of time has not been specified, shall be limited as follows:- A maximum of two and a half hours, with not more than twenty (20) minutes for the Mover in moving and five (5) minutes for any other Member speaking, including the Leader of the Majority Party and the Leader of the Minority Party and

the Chairperson of the relevant Committee (if the Committee Report is not moved by the Chairperson of the relevant Committee), and that ten (10) minutes before the expiry of the time, the Mover shall be called upon to reply; and further that priority in speaking shall be accorded to the Leader of the Majority Party and the Leader of the Minority Party, in that order.

Limitation of Debate on Audit Committee Reports

- V.** **THAT**, each speech in debate on **Reports of Audit Committees** be limited as follows: A maximum of sixty (60) minutes for the Mover in moving and thirty (30) minutes in replying, and a maximum of ten (10) minutes for any other Member speaking, except the Leader of the Majority Party and the Leader of the Minority Party, who shall be limited to a maximum of fifteen (15) minutes each; and that priority be accorded to the Leader of the Majority Party and the Leader of the Minority Party, in that order.



NOTICE PAPER

Tentative business for **Tuesday, June 11, 2024**

(Published pursuant to Standing Order 38(1))

It is notified that the following business is *tentatively* scheduled to appear in the Order Paper for Tuesday, June 11, 2024 –

A. COMMITTEE OF THE WHOLE HOUSE

The Kenya Drugs Authority Bill (National Assembly Bill No. 54 of 2022)
(The Hon. (Dr.) Robert Pukose, M.P)

(If not concluded on Thursday, June 06, 2024 – Afternoon Sitting)

B. THE STATUTE LAW (MISCELLANEOUS AMENDMENTS) BILL
(NATIONAL ASSEMBLY BILL NO. 67 OF 2023)

(The Leader of the Majority Party)

Second Reading

(If not concluded on Thursday, June 06, 2024– Afternoon Sitting)

C. THE MUNG BEANS BILL (SENATE BILL NO. 13 OF 2022)

(The Hon. Paul Nzengu, M.P. – Co-sponsor)

Second Reading

D. THE CARE AND PROTECTION OF CHILD PARENTS BILL (SENATE
BILL NO. 29 OF 2023)

(The Hon. Beatrice Kemei, M.P. – Co-sponsor)

Second Reading

E. THE LAND LAWS (AMENDMENT) (No. 2) BILL (NATIONAL
ASSEMBLY BILL NO. 76 OF 2023)

(The Leader of the Majority Party)

Second Reading

(If not concluded on Thursday, June 06, 2024 – Afternoon Sitting)

F. MOTION – **REPORTS OF THE AUDITOR-GENERAL ON THE**
NATIONAL GOVERNMENT CONSTITUENCIES
DEVELOPMENT FUND FOR NINE CONSTITUENCIES
IN BUNGOMA COUNTY

(The Chairperson, Decentralized Funds Accounts Committee)

(If not concluded on Thursday, June 06, 2024 - Afternoon Sitting)

- G. MOTION – PUBLIC PETITION ON FUNDS SPENT CONTRARY TO THE PROVISIONS OF ARTICLE 223 OF THE CONSTITUTION**
(The Chairperson, Public Petitions Committee)
(If not concluded on Thursday, June 06, 2024 – Afternoon Sitting)
- H. MOTION – REPORTS OF THE AUDITOR-GENERAL ON THE FINANCIAL STATEMENTS OF SPECIFIED STATE CORPORATIONS**
(The Chairperson, Public Investments Committee on Social Services, Administration and Agriculture)
(If not concluded on Thursday, June 06, 2024 – Afternoon Sitting)
- I. MOTION – ENHANCING REPORTING OF PARLIAMENTARY BUSINESS ON ONLINE PLATFORMS**
(The Chairperson, Committee on Parliamentary Broadcasting and Library)
(If not concluded on Thursday, June 06, 2024 – Afternoon Sitting)
- J. MOTION – INSPECTION VISIT TO SEMI-AUTONOMOUS INSTITUTIONS OF THE EAST AFRICAN COMMUNITY IN UGANDA**
(The Chairperson, Select Committee on Regional Integration)
(If not concluded on Thursday, June 06, 2024 – Afternoon Sitting)
- K. MOTION – IMPLEMENTATION STATUS OF REPORTS ON PETITIONS AND RESOLUTIONS PASSED BY THE HOUSE**
(The Chairperson, Select Committee on Implementation)
(If not concluded on Thursday, June 06, 2024 – Afternoon Sitting)
- L. MOTION – ALLEGED UNFAIR TRADE PRACTICES BY FOREIGN INVESTORS IN KENYA**
(The Chairperson, Departmental Committee on Trade, Industry and Cooperatives)
(If not concluded on Thursday, June 06, 2024 – Afternoon Sitting)
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APPENDIX

NOTICE OF PETITIONS, QUESTIONS & STATEMENTS

ORDER NO. 7 - STATEMENTS

It is **notified** that, pursuant to the provisions of Standing Order 44(2)(c), the following Statements will be **requested**—

No.	Subject	Member	Relevant Committee
1.	Power outage in Garissa County	<i>Hon. Umulker Harun, MP (Nominated Member)</i>	Energy
2.	Measures for mitigating the high cost of living	<i>Hon. (Bishop) Titus Khamala, MP (Lurambi)</i>	Finance and National Planning
3.	Collapsed <i>Athi River Bridge</i> in <i>Donyo Sabuk</i>	<i>Hon. Stephen Mule, MP (Matungulu)</i>	Transport and Infrastructure
4.	Regulation of Wildlife in Private Conservancies	<i>Hon. Mwangi Kiunjuri, MP (Laikipia East)</i>	Tourism and Wildlife
