| **Name of Department** | **Subject of Announcement** | **Update Periods** | **Related SOP/Circular or Legal Document Number/Year** |
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| Department of Pharmacovigilance and Controlled Substances  | Drug safety monitoring form | Whenever necessary | FVK-SOP-08 Procedure of Risk Management Plan Assessment /14.11.2019 |
| Department of Pharmacovigilance and Controlled Substances | Dear Doctor Letter | Whenever necessary | FVK-SOP-06 Procedure of Letter to Health Professionals/14.11.2019 |
| Department of Pharmacovigilance and Controlled Substances | Risk management materials  | Whenever necessary | FVK-SOP-08 Procedure of Risk Management Plan Assessment /14.11.2019 |
| Department of Pharmacovigilance and Controlled Substances | Announcement for Unions and Associations | Whenever necessary | FVK-SOP-04 Procedure of Safety Alert Assessment/20.04.2019 |
| Department of Pharmacovigilance and Controlled Substances | Pharmacovigilance contact points list | 1 month | SOP for Pharmacivigilance activities that will be carried in Provinces |
| Department of Rational Use of Medicines | Health CodingReference Server (HCRS) E-Prescription Medicines and Other Pharmaceutical Products List | Once a week | AİK-SOP03 Procedure of HCRS e-Prescription Medicines and OtherPharmaceutical Products List Creating andUpdating |
| Herbal and Supportive Products’ Department | Updating Medicinal Plant List  | When a new monograph of traditional herbal medicinal product being approved.  | BDU-SOP-09 Operating Procedure For Traditional Herbal and Homeopathic Medicinal Products Unit / 31.12.2019 |
| Department of Marketing Authorisation | FAQ | Whenever necessary | Coordination Unit Working Procedure/ İRD-SOP-14 / 31.12.2019 |
| Department of Marketing Authorisation | Announcements about Legislation/Legislation Changes | Whenever necessary | Coordination Unit Working Procedure / İRD-SOP-14 / 31.12.2019 |
| Department of Marketing Authorisation | List of API | Every month | CTD Pre Assessment Unit/ İRD-SOP-02 |
| Department of Marketing Authorisation | List of Marketing Authorised Products | Every week | Priority Evaluation Unit /IRD-SOP-01/27.09.2022 |
| Department of Marketing Authorisation | List of Medicines of which MA is cancelled  | First week of every month | IRD-SOP-11 Authorised Medicines Working Procedure/ 27.08.2022 |
| Department of Marketing Authorisation | SmPC and PILs  | Whenever necessary | IRD-SOP-08 Pharmacological Assessment Unit Working Procedure/27.09.2022 |
| Vice Prensidency of Inspectorate | Contact Information of Inspectors  | When there is a personnel change | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | Phase 1 Clinical Research Centers Permitted by Agency | 6 months | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | BA/BE Study Centers Permitted Agency | 6 months | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | Licensed Pharmaceutical Warehouses List | Continuous updates are made through the online system. | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | GMP Inspected Overseas Facilities and activities  | 6 months | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | Domestic Production Facilities Permitted by the Agency | Continuous updates are made through the online system. | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | Contact Information of Product Auditors | When there is a personnel change | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | List of Domestic Facilities Performed Routine Inspection | 6 months | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | List of Domestic Facilities to be Performed 6-Month Routine Inspections | 6 months | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | GDP Audit List | 6 months | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | GDP Risk Based Audit Plan | 6 months | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | List of Contracted Pharmacovigilance Service Organizations | 6 months | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | Unit Performances (with Administrative Activity Report) | 12 months | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | Announcements to Stakeholders | It has no period. Whenever necessary. | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | Revised Guidelines and Legislations | It has no period. Whenever necessary. | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | Announcements on the Sale Blockage of Pharmaceuticals | It has no period. Whenever necessary. | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |
| Department of Medicines Inspection | Announcements Regarding Recalls (Class I Recalls) | It has no period. Whenever necessary. | Communication Cooperation And Information Sharing GuidelineDHBY-SOP-23 |