Convenience Kits Interim Regulatory Guidance

This document is intended to provide guidance regarding a new premarket notification regulatory strategy for convenience kits. It does not create or confer any rights for or on any person and does not operate to bind FDA or the public. An alternative approach may be used if such approach satisfies the requirements of the applicable statute, regulations, or both. Public input on this document was not obtained prior to implementation because it is believed that this guidance presents a less burdensome policy that is consistent with public health.

Premarket Notification (510(k)) Staff
Program Operations Staff
Office of Device Evaluation

May 20, 1997

While this guidance document represents a final document, comments and suggestions may be submitted at any time for Agency consideration by contacting the Premarket Notification (510(k)) Section at 301-796-5640. For questions regarding the use or interpretation of this guidance, contact the Premarket Notification (510(k)) Section at 301-796-5640.

U.S DEPARTMENT OF HEALTH AND HUMAN SERVICES Food and Drug Administration Center for Devices and Radiological Health

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As part of the Center for Devices and Radiological Health's organizational transformation initiative, the 510(k) Process Reengineering Team has been carefully examining the Premarket Notification (510(k)) Program to insure that it is operating at peak efficiency and that the optimum public health benefit is achieved given the allotted resources.

During the preliminary examination of the requirements and execution of the 510(k) Program, the Team identified a change to the Program that may reduce the regulatory burden for assemblers of convenience kits. Outlined below is a new regulatory approach to be applied to convenience kits which could result in a decrease in the number of 510(k) submissions for these devices and, in so doing, will save ODE review resources. It is estimated that the Office of Device Evaluation (ODE) receives approximately 300 510(k) submissions per year that could be impacted by this regulatory approach.

Background

Under the current regulations, first time marketers of devices must submit a premarket notification and obtain clearance for a device before it can be lawfully introduced into interstate commerce. Assemblers/manufacturers of convenience kits who intend to market these devices for the first time are subject to these requirements.

Many of the kits that have been the subject of 510(k) review are comprised of legally marketed devices that are simply being assembled in kit form strictly for the "convenience" of the purchaser or user, thus the term "convenience kit." During the review of convenience kit 510(k)s, ODE has traditionally focused its efforts on the intended use(s) of the particular type of kit, the device components that are included in the kit, and the impact that any further processing may have on the kit and/or its components. The labeling for these kits tends to be rather basic, identifying the assembler or manufacturer of the particular kit and the intended use(s). Additional processing typically involves affixing the label to the outside packaging of the kit and terminal sterilization of the kit and its components.

New Regulatory Approach

Based on this experience, FDA believes that under certain circumstances premarket clearance for convenience kits may not be necessary to ensure protection of the public health. Accordingly, FDA intends to propose rulemaking to exempt certain, specifically identified convenience kits from the requirement of premarket notification. Until such a rule is in effect, FDA intends to exercise enforcement discretion regarding the requirement for premarket clearance for

convenience kits that conform with the specific limitations regarding intended use, components, and processing described below.

- 1. Intended Use Using the 510(k) database, ODE has developed the attached list of generic types of kits that the Agency believes represents established intended use(s) for convenience kits and that does not modify the intended use(s) of the individual kit components. FDA recognizes that additional types of kits may currently exist that should be subject to this guidance. Likewise, FDA anticipates that other types of kits may become eligible for consideration in time. Thus, the list may be periodically updated based on the Agency's experience with this new regulatory approach as well as on comments received from interested parties.
- 2. Components Convenience kits subject to this guidance should only include components that are either: (1) legally marketed preamendments devices, (2) exempt from premarket notification, or (3) have been found to be substantially equivalent through the premarket notification process. The components should be purchased in finished form, i.e., they should be packaged, labeled, etc., consistent with their legal marketing authorization.
- 3. Processing This guidance only applies in instances where the kit assembler/manufacturer concludes that further processing, if any, of the kit does not significantly affect the safety or effectiveness of any of the kit's components. In making this determination, FDA recommends that kit manufacturers follow the procedures described in FDA's guidance document entitled, "Deciding When to Submit a 510(k) for a Change to an Existing Device." It is the responsibility of each kit assembler/manufacturer to carefully consider the impact that any processing may have on the kit's components.

In the past, sterilization has been a focus of the 510(k) review for convenience kits; therefore, kit assemblers should carefully consider the impact of the sterilization process on individual kit components. If the kit's components may be sensitive to further processing, e.g., surgical sutures, the assemblers/manufacturers should take measures necessary to ensure that the components are not adversely affected by the reprocessing procedures.

In summary, FDA intends to exercise its enforcement discretion, i.e. not require 510(k) clearance, for convenience kits of a type matching one of those included on the attached list, consisting of components that have been cleared through the 510(k) process, and where the assembler/manufacturer is able to reasonably conclude that any further processing of the kit and its components does not significantly affect the safety or effectiveness of any of its components. Documentation to support each of these determinations should be maintained in the assembler's/manufacturer's files in accordance with the Quality System regulation (21 CFR 820) and should be available for FDA review if needed. It should be noted that while FDA intends to exercise enforcement discretion with respect to premarket notification requirements, assemblers/manufacturers of convenience kits are still required to comply with other general controls including registration, listing, prohibition against misbranding, and good manufacturing practices. In the future, FDA intends to propose rulemaking to formally exempt these types of kits from the requirement of premarket notification.

Limitations

FDA does not intend to propose regulatory changes relating to drug requirements for convenience kits. For convenience kits that contain components subject to regulation as drugs, the assembler/manufacturer should contact the Center for Drug Evaluation and Research, Division of Prescription Drug Compliance & Surveillance, at (301) 594-0101 regarding premarket requirements for the drug components in the kit.

This guidance is not intended to apply to kit 510(k)s that are under the jurisdiction of the Center for Biologics Evaluation and Research. For information regarding kits reviewed by CBER, the assembler/manufacturer should contact the Office of Blood Research and Review, Division of Blood Applications, Biologics Devices Branch at (301) 827-3524.

Effective Date

This guidance is effective immediately.

Types of Convenience Kits

The attached list of convenience kits was developed based on FDA's database and is organized by medical speciality. For purposes of this document, the terms "kit," "set," and "tray" are considered synonymous. This list may be periodically updated based on the Agency's experience with this guidance as well as on comments received from interested parties.

Name of Kit/Tray/Set and Intended Use (if Needed)

Anesthesiology Devices

Airway Suction Kit

Anesthesia Breathing Circuit Kit (Adult & Pediatric)

Anesthesia Conduction Kit

Anesthesia Kit

Arterial Blood Sampling Kit

Blood Specimen Collection Kit (Excludes HIV Testing)

Brachial Plexus Anesthesia Kit

Caudal Anesthesia Kit

Continuous Brachial Plexus Block Tray

Custom Anesthesia Tray

Epidural Anesthesia Kit

Glossopharyngeal Anesthesia Kit

Humidifier - Nebulizer Kit

Laryngoscope Kit

Nasal Endotracheal Tube Holder Kit

Nerve Block Tray

Orotrachael Intubation Guide Kit

Oxygen Administration Kit

Regional Anesthesia Kit

Spinal Anesthesia Kit

Spinal Epidural Anesthesia Kit

Tracheal Suction Set

Tracheobronchial Suction Catheter Kit

Tracheostomy and Nasal Suctioning Kit

Tracheotomy Care Kit

Cardiovascular Devices

Angiography/Angioplasty Kit

Angioscopic Valvulotome Kit

Cardiac Catheterization Kit

Cardioplegia Solution Administration Kit

Cardiopulmonary Bypass Catheter Kit

Cardiopulmonary Resuscitation Aid Kit

Cardiovascular Catheter Sheath Introducer Kit

Cardiovascular Procedure Kit

Cardiovascular Surgical Instruments Tray

Catheter Balloon Repair Kit

Catheter Guide Wire Kit

Catheter Introducer Kit

Central Venous Blood Pressure Kit

Central Venous Catheter Tray

CT Biopsy Tray

Digital Angiography Tray

Endothelial Cell Harvesting Kit

Laser Blood Flow Kit

Lead Introducer Kit

Percutaneous Atrial Catheter Kit

Percutaneous Sheath Introducer Kit

Phlebotomy Blood Collection Kit

Winged Intravenous Catheterization Kit

Clinical Chemistry and Clinical Toxicology Devices

Blood Alcohol Kit (Excludes HIV testing)

Blood and Urine Collection Kit (Excludes HIV testing)

Neonatal Blood Collection Kit and Screening Form (Excludes HIV testing)

Sex Crimes/Sexual Assault/Suspect Evidence Collection Kit (Excludes HIV testing)

Urine Collection Kit (Excludes HIV testing)

Urine Transport Kit (Excludes HIV testing)

Dental Devices

Dental Hygiene Kit
Dental Implant Surgical Tray
Dental Prophylaxis Kit
Denture Repair Kit
Endodontic Kit
Fixture Mount Kit
General Purpose Dental Tray
Gingival Retraction Kit
Lapping Tool Kit
Oral Surgery Tray
Plaque Disclosing Kit
Prophylaxis Kit
Restorative Instrument and Component Tray

Ear, Nose, and Throat Devices

Cricothyrotomy Kit
Ear Irrigation Kit
Ear, Nose, and Throat Surgical Tray
Myringotomy Procedure Kit
Oral Irrigation Kit
Pierced Ear/Entry Cleaning Tray
Tracheostomy Cleaning Tray
Tracheostomy Kit

Gastroenterology-Urology Devices

Anti-Regurgitation Blood Tubing Kit
Balloon Gastrotomy Tube Kit
Barium Enema Kit
Biopsy Needle Kit
Bladder Irrigation Kit
Body Fluids Barrier Kit
Bulb Irrigation Tray
Catheter Care Tray
Catheter (Insertion) Tray
Cholangiogram Catheter Kit
Continuous Arteriovenous Hemofiltration Catheter Kit
Declotting Tray

Dialysis Administration Kit

Dialysis On/Off Kit

Dialyzer Holder Set

Dual Lumen Catheter Tray

Electrosurgical Electrode Kit

Endoscope Introducer Kit

Endoscopic Cholangiogram Kit

Enema Kit

Enteral Administration Kit (Adult & Pediatric)

Esophageal Dilator Kit

Feeding Tube Kit

Femoral Catheter Kit

Flexible and Rigid Choledochoscope Kit

Foley Catheter Kit (Excludes HIV Testing)

Gastric Colonic Catheter Irrigation Kit

Gastric Irrigation and Aspiration Kit

Gastric Lavage Kit (Adult or Pediatric Use)

Gastro-Urology Biopsy Kit

Gastrointestinal Tube Kit

Gastrostomy Guide Wire Placement Kit

Gastrostomy Jejunostomy Feeding Tube Kit

Gastrostomy Tube Kit

Hemodialysis Tray

Kidney Perfusion Kit

Laparoscopic Cholangiogram Catheter Kit

Laparoscopic Gastrostomy Kit

Laparoscopic Jejunostomy Kit

Laparoscopy Kit

Male External Catheterization Kit (Excludes HIV Testing)

Multi-Lumen Hemodialysis Catheterization Kit

Nasogastric Feed Tube Kit

Nasogastric-Gastrojejunal Kit

Needle Catheter Jejunostomy Kit

Nephroscope Set

Non-Balloon Replacement Gastrostomy System

Ostomy Care Kit

Percutaneous Endoscopic Gastrostomy Kit

Percutaneous Replacement Gastrostomy Tube Kit

Percutaneous Cholangiogram Kit

Perineal Irrigation Kit

Peritoneal Dialysis Administration Kit

Peritoneal Dialysis Tubing Kit

Personal Erection Kit

Rectal Dilator Kit Replacement Gastrostomy Feeding Catheter Tray Single Needle with Uni-Directional Pump Kit Single Needle (Co-Axial Flow) Dialysis Kit Sterile Infant Gavage Kit Subclavian Catheter Kit Suprapubic Cystostomy Catheter Kit Universal Drainage Tray Urinary Irrigation Kit Urinary Drainage Collection Kit Urinary Catheterization Kit (Excludes HIV Testing)

General and Plastic Surgery Devices

Aspiration Tray Basic Set-Up Tray Bone Marrow Biopsy Tray **Bowel Kit**

Breast Biopsy/Localization Tray

Burn Dressing Tray

Burn Kit

Cement and Bone Removal Kit

Cholecystectomy Kit **Debridement Tray**

Decontamination Kit

Dressing Change Tray

First Aid Kit

Fluid Drainage Tray

General Purpose Instrument Tray

General Surgery Tray Herniorrhaphy Kit

Hysterectomy Kit

Incision and Drainage Tray Irrigation Kit (Wounds)

Laceration Tray

Laparoscopic Cholecystectomy Kit

Laparoscopic Sphincterotomy Kit

Liver Biopsy Kit

Minor Surgical Procedures Tray

Operating Room Accessories Table Tray

Paracentesis Tray

Pelviscopy Kit

Plastic Surgery and Accessories Kit

Shave Prep Kit

Single Use Instrument Tray

Skin Prep Tray

Staple Removal Kit

Sterile Puncture Tray

Sterilization Wrap Tray

Suction Catheter Kit

Surgical Instrument Tray

Surgical Retractor Kit

Suture Kit

Suture Removal Kit

Thoracic Catheter Insertion Tray

Tourniquet Kit

Ulcer Wound Tray

Vacuum Powered Body Fluid Collection Kit

Venous Ulcer Kit

Wound Drainage Kit

Wound Dressing Kit

General Hospital and Personal Use Devices

Admission Kit (Patient Utensil)

Anaphylactic Emergency Kit

Baby Care Kit

Blood Administration Kit

Blood Borne Pathogen Response Kit

Blood Sampling Kit (Excludes HIV Testing)

Blood Transfusion Kit

Body Fluid Barrier Kit

Body Fluid Clean Up Kit

Body Fluid Disposal Kit

Bone Marrow Collection/Transfusion Kit

Buret Administration Intravenous Kit

Catheter Repair Kit

Central Venous Catheter Dressing Change Kit

Chemotherapy Administration Kit

Chemotherapy Spill Clean-up Kit

Chest Drainage Kit

Clean-Up Kit

Delivery Room Apparel Kit

Emergency Response Safety Kit

Fever Monitoring Kit

I.V. Start Kit

Insect Sting Emergency Kit

Intra-Arterial Administration Kit

Intravenous Extension Tubing Kit

Lumbar Puncture Tray (Adult & Pediatric)

Medical Procedure Tray

Microsurgical Instrument Tray

Mid-Stream Collection Kit

Mortician's Kit

O.R. Scrub Prep Tray

Oral Administration Set

Parenteral Administration Kit

Parenteral Supply Kit

Percutaneous Introducer Kit

Peripheral Catheter Insertion Kit

Personal Protection Kit

Port Introducer Kit

Prep Kit

Pressure Monitoring (Air/Gas) Kit

Sitz Bath Kit

Snake Bite Kit

Snake Bite Suction Kit

Spill Kit

Survival Kit

Thermometer Kit

Thoracentesis Tray

Ultrasonic Aspiration Biopsy Kit

Umbilical Catheter Insertion Tray

Vascular Access Port Kit

Venipuncture Kit

Vomitus Clean-Up Kit

Neurological Devices

Endoscopic Shunt Placement Kit

Endoscopic Ventricular Catheter Placement Kit

Intracranial Pressure & Temperature Monitoring Kit

Lumbar Drainage Catheter Kit

Lymphangiographic Kit

Myelogram Kit

Neurological Test Kit

Neurological Tray

Obstetrical and Gynecological Devices

Amniocentesis Tray

Breast Pump Kit

Cervical Smear Kit

Cesarean Section Tray

Circumcision Tray

Culdocentesis Kit

Cytology Kit

D&C Tray

Delivery Kit

Emergency Obstetrical Kit

Endometrial Sampling Kit

Fetal Blood Sampling Kit (Excludes HIV Testing)

Forensic Evidence Sexual Assault Kit

Gynecological Laparoscopic Kit

Labor and Delivery Kit

Maternity Kit

Obstetrical Kit

Obstetrical Anesthesia Kit

Obstetrical Vacuum Delivery Kit

Pap Smear Kit

Paracervical Anesthesia Kit

Pelvic Exam Kit

Pudendal Anesthesia Kit

Seminal Fluid Collection Kit

Trocar Kit

Vaginal Examination Tray

Vasovasostomy Set

Opthalmic Devices

Eye Tray

IOL Tray

Phaco Kit

Surgical Eye Tray

Vitrectomy Kit

Orthopedic Devices

Disposable Joint Aspiration/Injection Kit
Endoscopic Plantar Kit
Graft Harvesting Kit
Laparoscopic Bone Anchor Urethropexy Instrument Kit
Orthopedic Tray
Pre-Drill Kit for Cartilage Re-surfacing
Surgical Cannulaes & Depth Gauge Kit for Arthroscopic Procedures
Surgical Kit
Transtibial Anterior Cruciate Ligament Kit

Physical Medicine Devices

Hand Accessory Kit

Radiology Devices

Arthrogram Tray
Biopsy Needle Guide Kit
Dental Tray
Discography Kit
MRI Disposable Kit
Prostate Seeding Kit
Radiographic Contrast Tray
Radiology-Diagnostic Kit