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## BioScience Products

To find out more about the BioScience products on the next few pages, as well as many others, please refer to our new Whatman BioScience Product Guide - # 12006B

### Nucleic Acid and Protein Sample Preparation

#### FTA® Card

Collect, archive, transport and purify nucleic acids, all at room temperature. Whatman FTA provides a remarkably easy way to collect and isolate nucleic acid samples for analysis. Simply apply virtually any type of biological sample to the FTA matrix and the nucleic acids are instantly captured and stabilized. Pathogens are inactivated, making samples safe to handle and ship. Store samples, including clones, at room temperature and analyze whenever you're ready.

#### FTA® Elute

The FTA Elute matrix is chemically treated with proprietary reagents that lyse cells upon contact causing the release of nucleic acids. DNA is recovered from the FTA Elute matrix through a simplified elution process using water and heat. Captured nucleic acid is easily released for multiple downstream applications in less than 30 minutes. FTA Elute Cards are stored at room temperature before and after sample application, reducing the need for laboratory freezers. FTA Elute rapidly inactivates organisms including blood-borne pathogens and eliminates the risk of contamination for the individuals handling the sample.

#### CloneSaver® Card

FTA Technology in a 96 Well format for high throughput applications. Designed for the collection, storage and purification of plasmid and BAC DNA from bacterial clones. DNA is stable at room temperature for at least 5 years (real-time data).

#### Elutrap®

The Elutrap System is designed to isolate nucleic acids and proteins from agarose or polyacrylamide gel slices by electroelution. Samples are concentrated in as little as 200 µl with excellent recoveries and without sample pretreatment or special buffers. Samples pass through a membrane which restricts the gel slice and are trapped by a molecular weight cutoff membrane for retention. The Elutrap System works with most horizontal gel electrophoresis chambers.



FTA



FTA Elute



CloneSaver Card



Elutrap

### Multiwell Plates

#### Protein Precipitation UNIFILTER® FF

Fast, easy and automatable protein precipitation. A fast, effective protein removal method for plasma and serum for high-throughput labs measuring drugs and metabolites. This high-quality filter plate replaces the lengthy centrifugation process with a vacuum filtration method, making sample preparation three times faster. It allows you to automate acetonitrile precipitation and speed up your research.



Protein Precipitation UNIFILTER FF

#### Protein Kinase Assay UNIFILTER®

Kinase assay in a 96 Well format. The Whatman Protein Kinase Assay filter plate incorporates a P81 filter in each well. P81 is a cation exchanger that binds peptides but does not bind unincorporated ATP, resulting in low non-specific background noise and high sensitivity in kinase assay.



Protein Kinase Assay UNIFILTER

#### ELISA UNIFILTER®

Better kinetics and simpler washing for ELISA. The Whatman ELISA plate allows researchers to utilize the excellent protein binding characteristics of nitrocellulose -49 - µg IgG per well in a 96 Well format. Solutions are easily vacuumed to waste using a vacuum manifold.



ELISA UNIFILTER

#### Phase Separation UNIFILTER®

Quick separation of halogenated solvents from an aqueous phase in a 96 Well format with no carryover and no close manual contact. Whatman 1PS media sealed into each well is a silicone treated media which remains impervious to aqueous solutions but organic solvents can go through.



Phase Separation UNIFILTER

#### Multi-Chem™ Microplates

Chemically resistant and low binding material microplate. Ideal for aggressive organic solvents such as DMF, TFA, THF, acetonitrile, chloroform and methylene chloride. Non-binding properties also make them ideal for storage of biological materials.

#### UNIPLATE™ 'V' Bottom Microplates

'V' bottom ensures maximum sample recovery. The 96 and 384 Well format UNIPLATE with 'V' bottom are ideal for applications with small sample volumes. The vertical sides of the well, combined with the 'V' design at the base of each well, ensure that all the material runs down the side walls and is channeled into the well base.

**Capmats**

Flexible capmats individually seal the top of each well. Capmats may be used on either filter or collection microplates.

**BugStopper® Microplate Capmat**

Sterile venting closures for 24 Well microplate cultures. 24 cultures (5-7 mL/sample) can be grown in a microplate, allowing easier handling than 24 test tubes. The autoclavable venting capmats significantly reduce evaporation rate and are perfect for extended growth of slow growing bacteria and fungi.

Protein Microarrays

**FAST Quant®**

FAST Quant kits are designed for high-throughput multiplex cytokine quantitation analyses. Each kit contains 64 arrays on FAST Slide surfaces with 8-10 monoclonal antibodies against a wide variety of cytokines per array, in triplicate. The most common cytokines for both human and mouse are represented in the FAST Quant system. The MicroSpot ELISA reaction is concentration dependent, making FAST Quant the fastest and most sensitive method of quantitating cytokines in a multiplex format.

**Serum Biomarker Chip**

The Serum Biomarker Chip allows proteomics researchers to pattern the molecular signature of human serum. The Serum Biomarker Chip is a single capture antibody array built on the FASTSlide dual pad platform. Each slide has an identical arrays of antibodies printed in triplicate. Two color fluorescent detection permits the comparison of the molecular signature of 120 human serum proteins between matched serum samples.

**Protein Array Services**

Several services are available for protein array researchers. Based on the FAST Quant System, the Quantitative Cytokine Array Processing & Data Analysis service will construct custom arrays from our antibody menu of 40 human and 19 murine specificities. Using the Serum Biomarker Chip service, researchers can send matched serum samples for analysis of 120 human serum proteins. Contract printing services are available to those researchers who wish to design their own protein array experiments. Whatman also offers a FAST Slide Scanning and Data Analysis service for smaller laboratories who do not wish to invest in instrumentation but want the value protein array experiments can bring them. Scientists at Whatman can also discuss and design entire protein array experiments from start to finish for those researchers just beginning protein array work.



Multi-Chem Microplates



BugStopper Microplate Capmat



FAST Quant



Serum Biomarker Chip



Protein Array Services

Blotting Products

**Protran®**

Protran nitrocellulose membranes are the most frequently specified transfer media in the world for a wide range of applications. Protran is made with 100% pure nitrocellulose for high binding capacities and low background. Protran is compatible with a variety of detection methods, including isotopic, chemiluminescent (luminol-based), colorimetric and fluorescent. Protran is wetted with an aqueous buffer which is ideal for proteins in aqueous environments. Protran is available in pore sizes of 0.1 µm, 0.2 µm and 0.45 µm for a wide variety of molecules.

**Minifold® I**

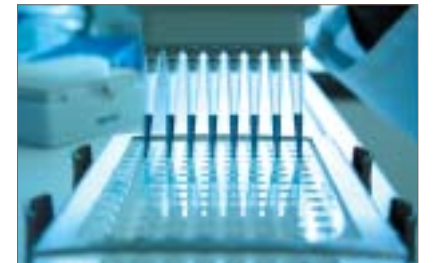
The Minifold System is for dot-, spot- or slot-blot arrays. The dot-, spot- or slot-blot plates are interchangeable on the vacuum manifold base, making the Minifold I System versatile for DNA or protein arrays. The spot- and dot-blot are in a 96-well format and the slot-blot has a 48-well format ideal for densitometric scanning. The Minifold I System is used with Protran, Optitran or Nytran membranes for blotting applications.

**TurboBlotter™**

The TurboBlotter is a rapid downward blotting device for the high-resolution transfer of DNA and RNA from agarose gels to blotting membranes. The traditional transfer setup has been turned upside down; no heavy weights are required for transfer. Alkaline DNA transfers can be performed in as little as 1 hour while neutral (SSC) transfers of DNA or RNA take only 3 hours. Complete kits have components for 5 transfers and replacement transfer packs are available.



Protran



Minifold I



TurboBlotter

# Filtration Simplified

## Basic Filtration Concepts and Terms

Selecting a filter with the appropriate properties can help you achieve accurate results and reach discovery faster. But with so many types of filters to choose from, how can you be sure you're making the right choice? Whatman has assembled this compilation of basic filtration concepts and terms to clarify the various options available to you and speed the process of selection.

### Airborne Particle Retention

Retention mechanisms for removing particulates from air or gas enable much higher efficiencies to be realized than those applicable to liquids. Efficiencies for air filtration are normally expressed as percent penetration or retention for a stated airborne particle size. In the United States, the Dioctyl Phthalate (DOP) test is commonly used wherein the filter is challenged with an aerosol containing 0.3  $\mu\text{m}$  particles.

### Ash Content

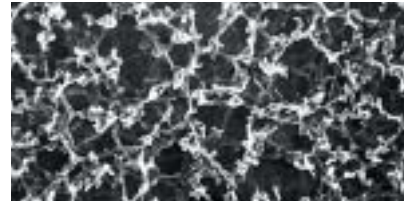
Determined by ignition of the cellulose filter at 900°C in air. Ash content is essential in gravimetric applications and also a useful measure of the level of general purity.

### Chemical Compatibility

It is very important to ensure that the pore structure of the filter media will not be impaired by exposure to certain chemicals. In addition, exposure to these chemicals should not cause the filter to shed fibers or particles, or add extractibles. Length of time exposure, temperature, concentration and applied pressure can all effect compatibility. Whatman has provided chemical compatibility charts to aid your membrane selection (see page 260).

### Depth Filters

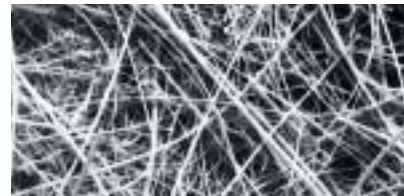
Depth filters are usually characterized as those which retain particles on the surface and within the filter matrix. All conventional fibrous filters (whether manufactured from cellulose, borosilicate glass microfiber or other fibrous material) are depth filters and are normally characterized by exhibiting good loading capacity.



*Membrane filters allow the efficient retention of submicron particulates and organisms.*



*Whatman cellulose filter papers exhibit particle retention levels down to 2.5  $\mu\text{m}$ .*



*Glass microfiber filters are manufactured by Whatman from 100% borosilicate glass.*



*Multigrade GMF 150 combines two filters in one for fast, effective multilayered filtration.*

### Herzberg Method

Whatman quantifies liquid flow rate for its range of filters by using a Herzberg flow rate test. Prefiltered deaerated water is applied to the test filter (effective area 10  $\text{cm}^2$ ) at a constant hydrostatic head (10 cm). The rate of the flow is measured in seconds per 100 mL. Flow rate can also be measured by the modified ASTM method which uses a quadrant folded filter held in a wire loop. It is not considered to be as reliable or consistent as the Herzberg test.

### Hydrophilic

Because hydrophilic filters possess an affinity for water and can be wetted with virtually any liquid, they are typically used for aqueous solutions.

### Hydrophobic

These types of filters repel water, and are thus best suited for venting or gas filtration applications.

### Liquid Flow Rate

Under practical filtration conditions, the liquid flow rate will depend on a number of factors, many of which will be specific to the solid/liquid being filtered. In order to compare filter performances, a standardized set of conditions is required which will characterize liquid flow rate for a given filter without the complicating secondary effects derived from the presence of particulates. Liquid flow rate is tested with prefiltered, deaerated water using a flat filter subjected to a constant hydrostatic head. Test methods based on quadrant folded filters are considered unreliable.

### Loading Capacity

This relates to the ability of a filter to load particulates into the fibrous matrix while maintaining a practical filtration speed and a workable pressure differential across the filter. In general, glass microfiber filters have a high loading capacity when compared with cellulose filters of the same retention rating and thickness. Membranes are inherently low in loading capacity. 'Choking life' is a measure of loading capacity.

### Particle Retention (Liquid)

In a filtration process, the particle retention efficiency of a depth-type filter is expressed in terms of the particle size (in  $\mu\text{m}$ ) at which a retention level of 98% of the total number of particles initially challenging the filter is obtained. It is customary to quote the retention levels at 98% efficiency to allow for secondary filtration effects. All Whatman depth filter grades have a published nominal retention rating determined on this basis.

### Pore Size

The pore size, usually stated in micrometers ( $\mu\text{m}$ ), of Whatman filter media is defined by the diameter of particles retained by the filter matrix. Pore size ratings, which can be either nominal or absolute, refer to the size of organisms or particles retained by the filter media.

### Prefilters

Prefilters are traditionally depth filters placed upstream from a membrane filter to significantly reduce the particulate loading in the system and thereby allow the membrane to operate efficiently at a light particulate loading.

### Screen or Surface Filters

Membrane filters are generally described as screen filters because particles are almost entirely trapped on the filter surface. The narrow effective pore size distribution of Whatman membrane filters is one of their major features.

## Filter Types and Filter Holders

### Filter Papers

Whatman qualitative and quantitative filter papers are, with few exceptions, manufactured from high-quality cotton linters which have been treated to achieve a minimum alpha cellulose content of 98%. These cellulose filter papers are used for general filtration and exhibit particle retention levels down to 2.5  $\mu\text{m}$ . There is a wide choice of retention/flow rate combinations to match numerous laboratory applications. The different groups of filter paper types offer increasing degrees of purity, hardness and chemical resistance. Whatman quantitative filter papers have extremely high purity for analytical and gravimetric work.

### Glass Microfiber Filters (GMF)

The unique properties of borosilicate glass microfibers enable Whatman to manufacture filters with retention levels extended into the submicron range. These depth filters combine fast flow rate with high loading capacity and retention of very fine particulates. Due to the high void volume exhibited by glass microfiber filters, the choking life is considerably extended beyond the life of a cellulose filter of similar retention. Whatman glass microfiber filters are manufactured from 100% borosilicate glass and most are completely binder-free. Binder-free glass microfiber filters will withstand temperatures up to 550°C and can therefore be used in gravimetric analysis where ignition is involved.

### Membrane Filters

Unlike cellulose and glass microfiber depth filters, membrane filters are conventionally classified as surface filters because the filter matrix acts as a screen and retains particulates almost entirely on the smooth membrane surface. The retention levels for these filters extend down to 0.02  $\mu\text{m}$  and allow the efficient retention of sub-micron particulates and organisms. Water microbiology and air pollution monitoring are major applications of membranes.

### Prefilters

The life of a membrane filter can be extended many times by placing a prefilter upstream of the membrane. The total particulate load challenging the membrane is considerably reduced thus allowing the membrane to operate efficiently.

### Standard Circle Funnel Volumes

The maximum practical volume of the most popular circle sizes (quadrant folded) is given in the following chart. Membrane and glass microfiber filters are used flat.

Diameter (cm)	Volume (mL)
9	15
11	20
12.5	35
15	75
18.5	135
24	300

### Types of Filter Holders

A filter matrix requires a suitable support structure to enable it to be used for the filtration of liquids or gases. One of the simplest forms of holder is the conical glass filter funnel into which a quadrant folded or fluted filter paper is placed (1). Some applications require additional motivating force for the solid particulate/liquid separation to occur (i.e., vacuum assisted filtration). This type of filtration can be carried out in a one-piece Büchner style funnel (2) where the filter is used flat on a perforated base sealed into the funnel. Due to the difficulties encountered in cleaning this type of funnel, the demountable 3-piece funnel was developed (3). The Whatman 3-Piece Filter Funnel is fully demountable and enables the filter paper to be securely clamped between the support plate and filter reservoir flange. Membrane holders (4) incorporate either sealed-in sintered glass or removable stainless steel mesh supports for the membrane. Syringe and in-line filters are also available. Large diameter membranes are typically used in pressure holders.

### Selecting the Right Filter

The selection of a laboratory filter depends on the conditions and objectives of the experiment or analytical procedure.

The three most important characteristics of any laboratory filter are:

- Particle retention efficiency
- Fluid flow rate through the filter
- Loading capacity

In addition, according to the particular application, other important characteristics may require examination. For instance, wet strength, chemical resistance, purity and ash level may assume equal importance under certain circumstances.



1



2



3



4

## Standard 58° or 60° Funnels

Glass/Polyethylene	
Funnel Diameter (mm)	Filter Paper Size (cm)
35	5.5
45	7.0
55	9.0
65	11.0
75	12.5
90	15.0
100	18.5
160	24.0
180	32.0
220	40.0
260	50.0

## Büchner Funnel Filter Selection

Diameter (mm)	Perforated Area (mm)	Filter Paper Size (mm)
43	32	42.5
63	42	55
83	60	75
100	77	90
114	95	110
126	105	125
151	135	150
186	160	185
253	213	240

## Typical Particle Sizes

Gelatinous Precipitates	μm
Metal Hydroxides	25–40
Precipitated Silica	25–40
Crystalline Precipitates	
Ammonium Phosphomolybdate	20
Calcium Oxalate	15
Lead Sulfate	10
Barium Sulfate (hot ppt.)	8
Barium Sulfate (cold ppt.)	3
Blood Cells	
Platelets	2–3
Erythrocytes (average)	7.0
Polymorphs	8–12
Small Lymphocytes	7–10
Large Lymphocytes	12–15
Monocytes	16–22
Bacteria*	
Cocci	0.5
Bacilli	1.0 x (1.0–1.0)
<i>Serratia marcescens</i>	0.5 x (0.5–1.0)
<i>Pneumococcus</i>	1.0
<i>Bacillus tuberculosis</i>	0.3 x (2.5–3.5)
Amoeba	12–30
<i>E. coli</i>	0.5 x (1.0–3.0)
Smallest Bacteria	0.22
Other Microorganisms, etc.	
Yeast Cells	2.0–8.0
Tobacco Smoke	0.5
Colloids	0.06–0.30
Rye Grass Pollen	34
Ragweed Pollen	20
Puffball Spores	3.3

\* Where bacteria are rod-shaped, range of lengths is given in brackets

## Product Selection

## Compatibility of Membranes

Solvent	ANP	CA	CN	PC	PE	GMF	NYL	PP	dpPP	PSU	PES	PTFE	PVDF
Acetic Acid, 5%	R	LR	R	R		R	R	R	R	R	R	R	R
Acetic Acid, Glacial	R	NR	NR			R	LR	R	R	R	R	R	R
Acetone	R	NR	NR	NR	R	R	R	R	R	NR	NR	R	NR
Acetonitrile	R	NR	NR			R	R	R	R	NR	R	R	R
Ammonia, 6N	NR	+	NR	NR	LR	LR	R	R	R	R	R	R	LR
Amyl Acetate	LR	NR	NR	R	R	R	R	R	R	NR	LR	R	LR
Amyl Alcohol	R	R	R			R	R	R	R	R	NR	R	R
Benzene*	R	R	R	LR	R	R	LR	LR	LR	NR	R	R	R
Benzyl Alcohol*	R	LR	LR	LR	R	R	LR	R	R	NR	NR	R	R
Boric Acid	R	R	R	R	R	R	LR	R	R	R	+	R	R
Butyl Alcohol	R	R	R	R	R	R	R	R	R	R	R	R	R
Butyl Chloride*		+				R	NR	NR	NR	+	+	R	R
Carbon Tetrachloride*	R	NR	R	LR	R	R	LR	LR	LR	NR	R	R	R
Chloroform*	R	NR	R	NR	R	R	NR	LR	LR	NR	NR	R	R
Cyclohexanone	R	NR	NR			R	NR	R	R	NR	NR	R	R
Chlorobenzene	R	+	R			R	+	+	+	+	NR	R	R
Citric Acid		+	+			R	LR	+	+	+	R	R	R
Cresol		NR	R			R	NR	R	R	NR	NR	R	NR
Cyclohexane	R	R	R	R	R	R	R	R	R	R	R	R	R
Diethyl Acetamide		R	NR			R	R	R	R	NR	+	R	NR
Dimethyl Formamide	LR	NR	NR			R	R	R	R	NR	NR	R	NR
Dioxane	R	NR	NR	NR	R	R	R	R	R	NR	LR	R	LR
DMSO	LR	NR	NR	NR	R	R	R	R	R	NR	NR	R	LR
Ethanol	R	R	NR	R	R	R	R	R	R	R	R	R	R
Ethers	R	LR	LR	R	R	R	R	R	R	R	R	R	LR
Ethyl Acetate	R	NR	NR	LR	R	R	R	R	R	NR	NR	R	LR
Ethylene Glycol	R	LR	LR	R	R	R	R	R	R	R	R	R	R
Formaldehyde	LR	LR	R	R	R	R	R	R	R	R	R	R	R
Freon TF	R	R	R	R	R	R	R	R	R	R	R	R	R
Formic Acid		LR	LR			R	NR	R	R	LR	R	R	R
Hydrochloric Acid, Conc	NR	NR	NR	R	NR	R	NR	LR	LR	R	R	R	R
Hydrofluoric Acid		NR	NR			NR	NR	LR	LR	+	+	R	R
Hexane	R	R	R	R	R	R	R	R	R	R	R	R	R
Isobutyl Alcohol	R	R	LR	R	R	R	R	R	R	R	+	R	R
Isopropyl Alcohol	R	R	LR			R	R	R	R	NR	+	R	R
Methanol	R	R	NR	R	R	R	R	R	R	R	R	R	R
Methyl Ethyl Ketone	R	LR	NR	LR	R	R	R	R	R	NR	NR	R	R

contd &gt;

Solvent	ANP	CA	CN	PC	PE	GMF	NYL	PP	dpPP	PSU	PES	PTFE	PVDF
Methylene Chloride*	R	NR	LR			R	NR	LR	LR	NR	NR	R	R
Nitric Acid, Conc		NR	NR	R	NR	R	NR	NR	NR	NR	NR	R	NR
Nitric Acid, GN		LR	LR			R	NR	LR	LR	LR	LR	R	LR
Nitrobenzene*	LR	NR	NR	NR	R	R	LR	R	R	LR	NR	R	R
Pentane	R	R	R	R	R	R	R	R	LR	R	R	R	R
Perchloro Ethylene	R	R	R			R	R	R	LR	NR	NR	R	R
Pyridine	R	NR	NR	NR	R	R	LR	R	R	NR	NR	R	R
Phenol 0.5%	LR	LR	R			R	R	R	R	NR	NR	R	R
Sodium Hydroxide, 6N	NR	NR	NR	NR	NR	NR	LR	R	R	R	R	R	NR
Sulfuric Acid, Conc	NR	NR	NR	NR	NR	R	NR	NR	R	NR	NR	R	NR
Tetrahydrofuran	R	NR	NR			R	R	LR	LR	NR	NR	R	R
Toluene*	R	LR	R	LR	R	R	LR	LR	LR	NR	NR	R	R
Trichloroethane*	R	NR	LR	NR	R	R	LR	R	R	NR	R	R	R
Trichloroethylene*	R	+	R			R	NR	R	R	NR	NR	R	R
Water	R	R	R	NR	R	R	R	R	R	R	R	NR	R
Xylene*	R	R	R			R	LR	LR	LR	NR	LR	R	R

R = Resistant; LR = Limited Resistance; NR = Not Recommended; + = Insufficient Data; \* = Short Term Resistance of Housing  
The above data is to be used as a guide only. Testing prior to application is recommended.

## Membrane Abbreviations:

ANP – Anopore  
CA – Cellulose Acetate  
CN – Cellulose Nitrate  
PC – Polycarbonate  
PE – Polyester  
GMF – Glass Microfiber  
NYL – Nylon  
PP – Polypropylene  
dpPP – Depth Polypropylene  
PSU – Polysulfone  
PES – Polyethersulfone  
PTFE – Teflon  
PVDF – Polyvinylidene Fluoride

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7060-2508	54	7195-002	37	UN203APEAQU	133
7060-2510	54	7195-004	37	UN203APENYL	133
7060-2511	54	7195-009	37	UN203APEORG	133
7060-2513	53	7402-001	44	UN203APEPES	133
7060-2514	54	7402-002	44	UN203APEPP	133
7060-4702	53	7402-004	44	UN203APUAQU	133
7060-4710	53	7402-009	44	UN203APUDPP	133
7060-4713	53	7402-009	44	UN203APUDPP	133
7060-4716	53	7404-001	44	UN203APUGMF	133
7061-2502	53	7404-002	44	UN203APUNYL	133
7061-2504	53	7404-004	44	UN203APUORG	133
7061-2510	53	7404-009	44	UN203APUPES	133
7061-4702	54	7408-004	44	UN203APUPP	133
7063-2502	58	7582-002	50	UN203NPEAQU	131
7063-2504	58	7582-004	50	UN203NPENYL	131
7063-4702	58	7585-004	50	UN203NPEORG	131
7091-4710	54	7590-004	50	UN203NPEPES	131
7141-004	37	7592-104	47	UN203NPEPP	131
7141-104	37	800195	55	UN203NPERC	132
7141-114	37	AV115NPEORG	128	UN203NPUAQU	131
7141-124	37	AV115NPUAQU	128	UN203NPUDPP	131
7141-154	37	AV115NPUNYL	128	UN203NPUGMF	131
7141-204	37	AV115NPUORG	128	UN203NPUNYL	131
7153-004	41	AV115UGMF	128	UN203NPUORG	131
7153-104	41	AV125EAQU	128	UN203NPUPES	131
7181-002	37	AV125ENAO	128	UN203NPUPP	131
7181-004	37	AV125EORG	128	UN203NPURC	132
7182-001	37	AV125EPP	128	UN503NPEAQU	131
7182-002	37	AV125NPUAQU	128	UN503NPENYL	131
7182-004	37	AV125NPUPSU	128	UN503NPEORG	131
7182-009	37	AV125SAQU	128	UN503NPEPES	131
7182-014	37	AV125SNAO	128	UN503NPEPP	132
7184-001	37	AV125SORG	128	UN503NPERC	132
7184-002	37	AV125UAQU	128	UN503NPUAQU	132
7184-003	37	AV125UCA	128	UN503NPUDPP	132
7184-004	37	AV125UGMF	128	UN503NPUGMF	132
7184-008	37	AV125UNAO	128	UN503NPUNYL	132
7184-009	37	AV125UORG	128	UN503NPUORG	132
7184-014	37	AV125UPP	128	UN503NPUPES	132
7184-029	37	AV125URCT	128	UN503NPUPP	132
7186-002	37	AV525BGMF	128	UN503NPURC	132
7186-004	37	AV525UAQU	128	US203NPEAQU	132
7187-114	41	AV525UNAO	128	US203NPENYL	132
7188-002	37	AV525UORG	128	US203NPEORG	132
7188-003	37	AVST25040	128	US203NPEPES	132
7188-004	37	CR0000006	133	US203NPEPP	132
7188-009	37	UN113EAQU	134	US203NPUPP	132
7190-002	37	UN113ENYL	134	US203NPUAQU	132
7190-004	37	UN113EORG	134	US203NPUDPP	132
		UN113UAQU	134	US203NPUGMF	132
				US203NPUNYL	132
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				US203NPUPES	132
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				US503NPEAQU	132
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				US503NPUGMF	132
				US503NPUNYL	132
				US503NPUORG	132
				US503NPUPES	132
				US503NPUPP	132

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