



import vendor manual 2.7

Big Lots Import Vendor Manual 2.7 APPENDIX – Table of Contents

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SHIPPING MARKS

CSC DISTRIBUTION, INC. - MONTGOMERY DC# 870

In order to avoid confusion and eliminate mistakes careful attention must be given to all carton markings. The following illustrates proper carton markings.

CARTON MARKINGS MUST BE IN BLACK!!!!

THE DEPARTMENT NUMBER IS THE DEPARTMENT NUMBER ON THE COMPUTER GENERATED PURCHASE ORDER.

Side Mark (both sides)	End Mark (both ends)	Inner Pack Mark (one end)
CSC DISTRIBUTION Montgomery, AL. PO# SKU# Dept # Carton of Country of Origin	SKU# Description Master Pack Inner Pack GW/NW (kgs.) Cube	SKU# Description Pack
Posts Sides	SKU# 310110000 BOTTLE OPENER Master Pack 288 Inner Pack 24 24 kgs/22kgs5 CBM	SKU# 310110000 BOTTLE OPENER 24 EA.
Both Sides	l Both Ends	One End Only

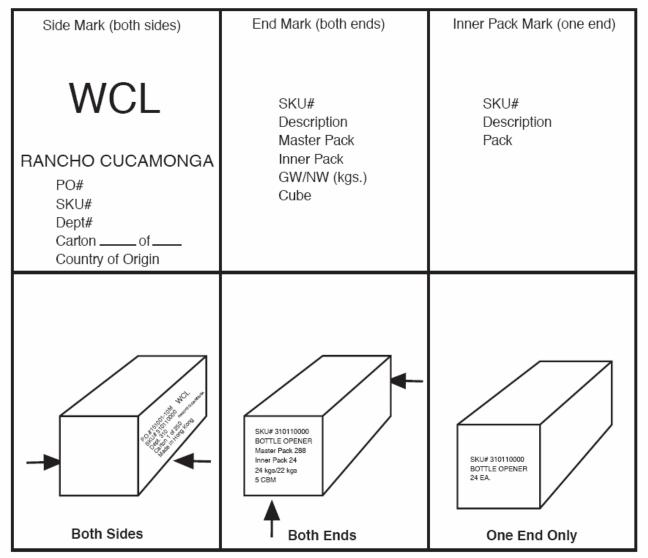
SHIPPING MARKS

WEST COAST LIQUIDATORS, INC. - RANCHO CUCAMONGA DC#873

In order to avoid confusion and eliminate mistakes careful attention must be given to all carton markings. The following illustrates proper carton markings.

CARTON MARKINGS MUST BE IN BLACK!!!!

THE DEPARTMENT NUMBER IS THE DEPARTMENT NUMBER ON THE COMPUTER GENERATED PURCHASE ORDER.



SHIPPING MARKS

DURANT DC, LLC - DURANT DC# 879

In order to avoid confusion and eliminate mistakes careful attention must be given to all carton markings. The following illustrates proper carton markings.

CARTON MARKINGS MUST BE IN BLACK!!!!

THE DEPARTMENT NUMBER IS THE DEPARTMENT NUMBER ON THE COMPUTER GENERATED PURCHASE ORDER.

Side Mark (both sides)	End Mark (both ends)	Inner Pack Mark (one end)	
DURANT DC Durant, OK PO# SKU# Dept # Carton of Country of Origin	Sku # Description Master Pack Inner Pack GW/NW (kgs.) Cube	Sku# Description Pack	
Poth Sidos	SKU# 310110000 BOTTLE OPENER Master Pack 288 Inner Pack 24 24 kgs/22kgs5 CBM	SKU# 310110000 BOTTLE OPENER 24 EA.	
Both Sides	Both Ends	One End Only	

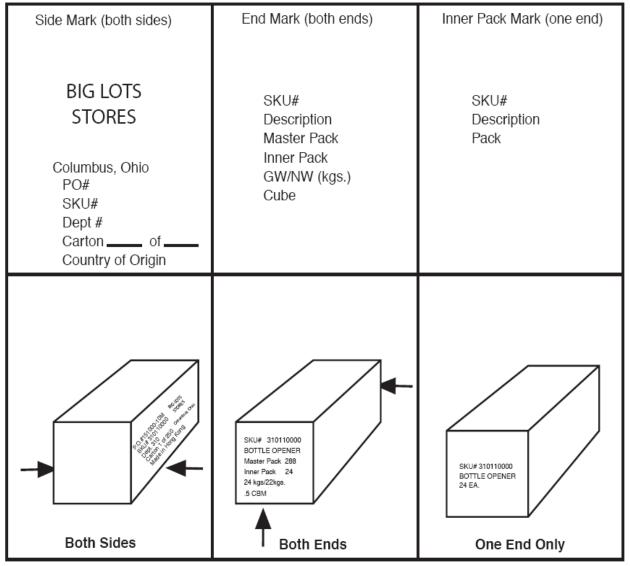
SHIPPING MARKS

BIG LOTS STORES, INC. - COLUMBUS DC# 890

In order to avoid confusion and eliminate mistakes careful attention must be given to all carton markings. The following illustrates proper carton markings.

CARTON MARKINGS MUST BE IN BLACK!!!!

THE DEPARTMENT NUMBER IS THE DEPARTMENT NUMBER ON THE COMPUTER GENERATED PURCHASE ORDER.



SHIPPING MARKS

BIG LOTS FURNITURE - REDLANDS DC# 893

In order to avoid confusion and eliminate mistakes careful attention must be given to all carton markings. The following illustrates proper carton markings.

CARTON MARKINGS MUST BE IN BLACK!!!!

THE DEPARTMENT NUMBER IS THE DEPARTMENT NUMBER ON THE COMPUTER GENERATED PURCHASE ORDER.

Side Mark (both sides)	End Mark (both ends)	Inner Pack Mark (one end)
BIG LOTS FURNITURE Redlands, CA PO# SKU# Dept # Carton of Country of Origin	SKU# Description Master Pack Inner Pack GW/NW (kgs.) Cube	SKU# Description Pack
Both Sides	SKU# 310110000 BOTTLE OPENER Master Pack 288 Inner Pack 24 24 kgs/22kgs5 CBM Both Ends	SKU# 310110000 BOTTLE OPENER 24 EA. One End Only

SHIPPING MARKS

WISCONSIN TOY - COLUMBUS DC# 964

In order to avoid confusion and eliminate mistakes careful attention must be given to all carton markings. The following illustrates proper carton markings.

CARTON MARKINGS MUST BE IN BLACK!!!!

THE DEPARTMENT NUMBER IS THE DEPARTMENT NUMBER ON THE COMPUTER GENERATED PURCHASE ORDER.

Side Mark (both sides)	End Mark (both ends)	Inner Pack Mark (one end)
WISCONSIN TOY Columbus, Ohio PO# Carton of Country of Origin	Item# Description Master Pack Inner Pack GW/NW (kgs.) Cube	Item# Description Pack
Both Sides	Item# A-3461 Description PLUG Master Pack 288 Inner Pack 24 GW/NW (kgs) 40/38 Cube .5 CBM Both Ends	Item# A 332 Description PLUG Pack 24 EA. One End Only

SHIPPING MARKS

CONSOLIDATED INTERNATIONAL - COLUMBUS DC #975

In order to avoid confusion and eliminate mistakes careful attention must be given to all carton markings. The following illustrates proper carton markings.

CARTON MARKINGS MUST BE IN BLACK!!!!

THE DEPARTMENT NUMBER IS THE DEPARTMENT NUMBER ON THE COMPUTER GENERATED PURCHASE ORDER.

Side Mark (both sides)	End Mark (both ends)	Inner Pack Mark (one end)
CONSOLIDATED INTERNATIONAL Columbus, Ohio PO# Carton of Country of Origin	Item# Description Master Pack Inner Pack GW/NW (kgs.) Cube	Item# Description Pack
	Item# A-3461 Description PLUG Master Pack 288 Inner Pack 24 GW/NW (kgs) 40/38 Cube .5 CBM	Item# A 332 Description PLUG Pack 24 EA.
Both Sides	Both Ends	One End Only

SHIPPING MARKS

CLOSEOUT DISTRIBUTION, INC. - TREMONT DC# 874

In order to avoid confusion and eliminate mistakes careful attention must be given to all carton markings. The following illustrates proper carton markings.

CARTON MARKINGS MUST BE IN BLACK!!!!

THE DEPARTMENT NUMBER IS THE DEPARTMENT NUMBER ON THE COMPUTER GENERATED PURCHASE ORDER.

Side Mark (both sides)	End Mark (both ends)	Inner Pack Mark (one end)	
CLOSEOUT DISTRIBUTION Tremont, Pa. PO# SKU# Dept # Carton of Country of Origin	SKU# Description Master Pack Inner Pack GW/NW (kgs.) Cube	SKU# Description Pack	
	SKU# 310110000 BOTTLE OPENER Master Pack 288 Inner Pack 24 24 kgs/22kgs5 CBM	SKU#310110000 BOTTLE OPENER 24 EA.	
Both Sides	Both Ends	One End Only	

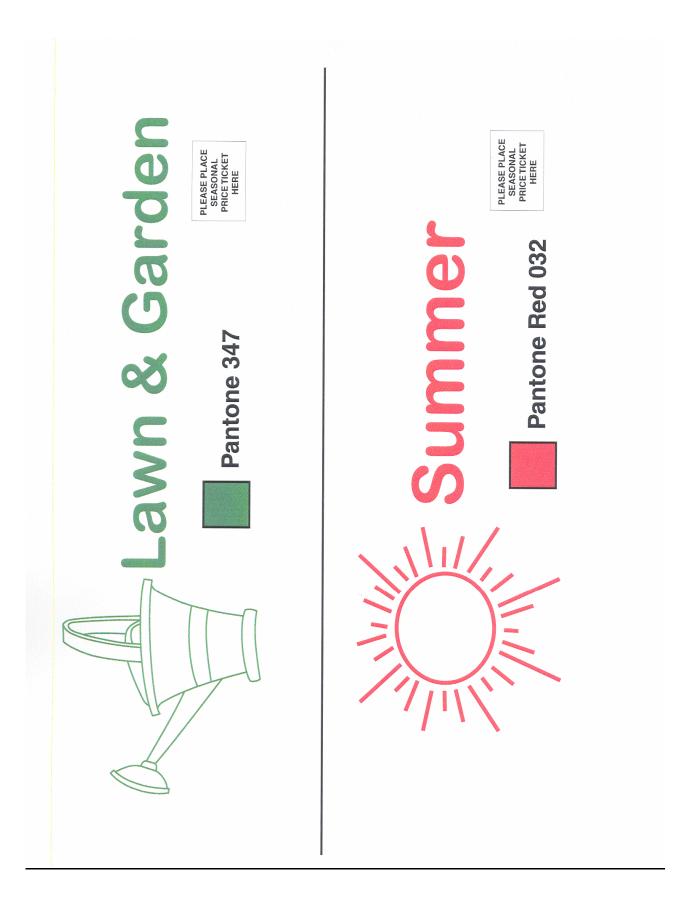
IMPORTANT: NEW SEASONAL CARTON MARKINGS

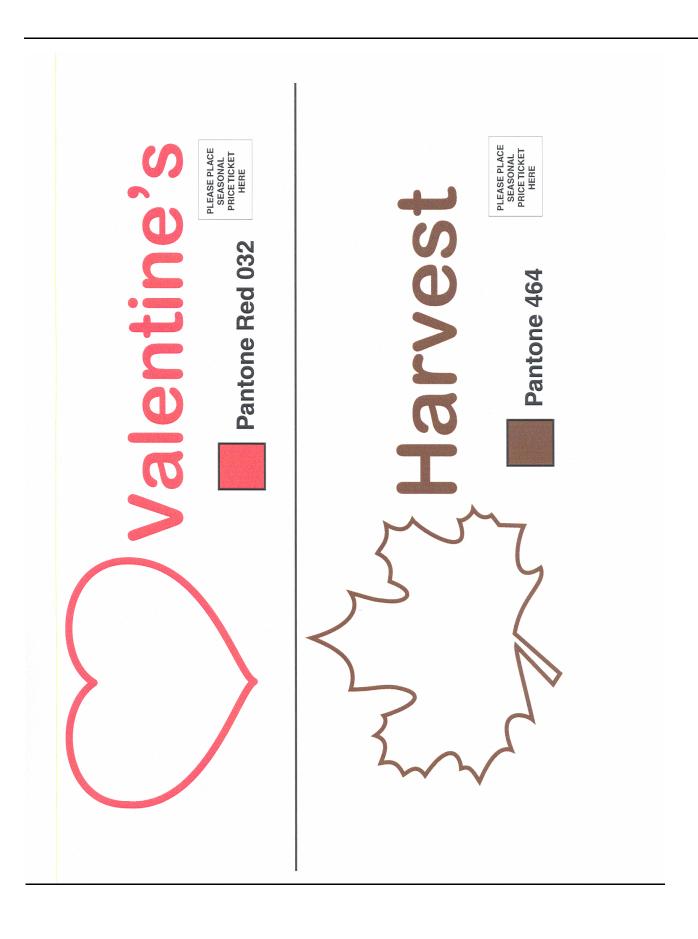
In order for our stores to more easily identify seasonal merchandise, we are adding the following seasonal specifics to the markings on our shipping cartons.

Print the seasonal carton markings in the upper left hand corner of the two main panels of the shipping carton. We have supplied graphics, (see the following pages), with horizontal orientation for seasonal carton markings. Please place the seasonal graphics so the width fills at least half of the carton panel. Match the seasonal department number to the department number on each purchase order.

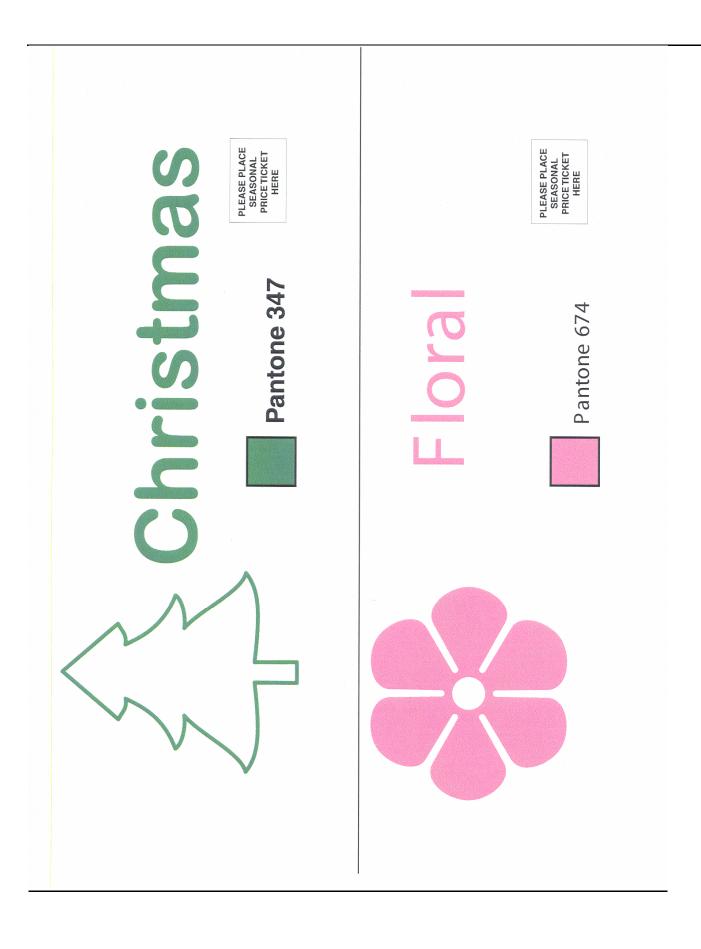
Be sure to print the seasonal carton markings in the correct pantone number shown on the following pages. This pantone number listed is for your reference only. Note that the rest of the carton markings must be in black.













Introduction

This document outlines the standard graphics and symbols to be used by both domestic and international vendors to mark containers shipped to Big Lots. Additionally, it serves as a guideline to Big Lots' Distribution & Transportation Services associates and Big Lots' Store associates for interpretation and use of container symbols.

Compliance with shipping container symbols throughout the supply chain is critical for proper and safe container handling and storage. Following these guidelines will help minimize the risk of product damage during transportation and handling. There are a few exceptions when these symbols can be disregarded during one or more stages of the supply chain. Such exceptions are on an individual basis and will be highlighted in this document. Additional requests for waivers or exceptions must be requested in writing from Big Lots.

The container symbols in this document are based on, and are consistent with, either the American Society for Testing and Materials (ASTM) standard D 5445 (Standard Practice for Pictorial Markings for Handling of Goods) or the International Standards Organization (ISO) standard ISO 780 Pictorial Marking for Handling of Goods. As these guidelines and standards are updated by the governing bodies, Big Lots will review the updates and refine the labeling standards accordingly.

The symbols in this document provide instructions for special handling of shipping containers. It is not necessary to apply all of the symbols on every container. Application of the shipping symbols is based on the shipping and handling requirements of the product/package system.

The size of each graphic or label should be determined according to the table below.

Dimension of shortest	Minimum Dimension of Individual Graphic or Label	
side panel in inches	Height (Inches) Width (Inches	
36 or greater	4	3
24 to 36	2 3/4	1 3/4
24 or less	2 1/4	1 1/2

Note: in this document the term shipping container, carton and package are used interchangeable.

Container graphics

1. Fragile - Handle with care



Figure 1: "Fragile – Handle with care" symbol

This symbol indicates that the contents of the container are fragile and that the container should be handled with care. This symbol should be used when the contents of the container contains glass, mirror or ceramic pots.

	Vendor	DC	Store
Must be complied with by:	X	X	X

Notes:

♦ This symbol should be placed on at least two adjacent sides of the container in the upper corner as shown below and in Pantone 032.

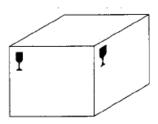


Figure 2: "Fragile – Handle with care" symbol location

♦ Corresponds with ISO 7000 / No. 0621.

2. This way up



Figure 3: "This way up" symbol

This symbol indicates the <u>required</u> upright position of the container. The symbol should only be used where consideration of the safety of the contents necessitates that the container be stored or stacked with the top surface up, and on cartons that should never be inverted or laid on their side.

	Vendor	DC	Store
Must be complied with by:		X	X

* As container cubing is a priority for Big Lots due to transportation costs, vendors are allowed to stack cartons in non-upright positions in order to maximize ocean container/trailer cubing as long as there is no major risk of product damage. Under no circumstances should the "This way up" arrows be facing down.

Notes:

- Arrows should never be shown on top of the package.
- ◆ This symbol shall be shown near the left hand upper corner on all four upright sides of the package and in Pantone 032.

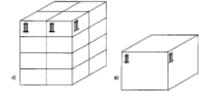


Figure 4: "This way up" symbol location

♦ When both the "Fragile – handled with care" and "This way up" symbols are used on a container, the latter should be closest to the corner as shown in the following figure.

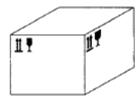


Figure 5: "This way up" and "Handle with care" symbol location

♦ Corresponds with ISO 7000 / No. 0623.

3. Stacking limitation by number

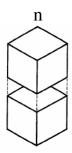


Figure 7: "Stacking limitation by number" symbol

This symbol indicates the maximum number of the same container (package) that can be stacked on top of <u>like</u> containers (packages), where 'n' is the maximum number of containers that can be stacked.

For example, if a container can only be stacked 5 high the "stacking limitation by number" symbol on the container should be:

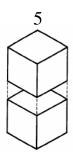


Figure 8: "Stacking limitation by number" symbol example

	Vendor	DC	Store
Must be complied with by:		X	X

^{*} As container cubing is a priority for Big Lots due to transportation costs, vendors are allowed to exceed the stacking limitation by number by a reasonable number in order to maximize ocean container/trailer cubing as long as there is no major risk of product damage.

Notes:

- ♦ The number of like containers that can be stacked on top of a package is typically determined by vendors by means of compression testing.
- ♦ The symbol should be located on at least 2 adjacent sides of the container and in Pantone 032.

4. Stacking limitation by weight

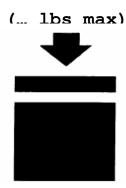


Figure 9: "Stacking limitation by weight" symbol

This symbol indicates the maximum weight of <u>unlike</u> packages that can be stacked on top a container. The maximum weight should be noted in pounds at the top of the symbol. For example, if no more than 80 pounds can be stacked on top of a container the "stacking limitation by weight" symbol on the package should be:

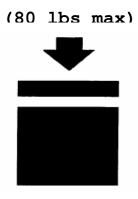


Figure 10: "Stacking limitation by weight" symbol example

	Vendor	DC	Store
Must be complied with by:		X	X

* As container cubing is a priority for Big Lots due to transportation costs, vendors are allowed to exceed the stacking limitation by weight by a reasonable amount in order to maximize ocean container/trailer cubing as long as there is no major risk of product damage.

Notes:

- ♦ The maximum weight of unlike packages that can be stacked on top of a container is typically determined by vendors by means of compression testing.
- ♦ This symbol should be located on at least 2 adjacent sides of the container and in Pantone 032.

♦ Corresponds with ISO 7000 / No. 0630.

5. Clamp here



Figure 15: "Clamp here" symbol

This symbol indicates that it is safe to clamp the container from the side where it is displayed and there is adequate packaging protecting the product from potential damaged caused by clamp trucks.

	Vendor	DC	Store
Must be complied with by:	X	X	X

Notes:

- ♦ This symbol should be located on two opposite sides of the container so it is in visible range of a clamp truck operator and in Pantone 032.
- ♦ Corresponds with ISO 7000 / No. 0631.

6. Do not clamp here



Figure 16: "Do not clamp here" symbol

This symbol indicates that the container should not be clamped from the side where the symbol is displayed as the packaging is not designed to withstand compression forces caused by clamp trucks.

	Vendor	DC	Store
Must be complied with by:	X	X	X

Notes:

♦ This symbol should be located on two opposite sides of the container so it is in visible range of a clamp truck operator and in Pantone 032.

7. Team lift

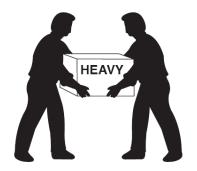


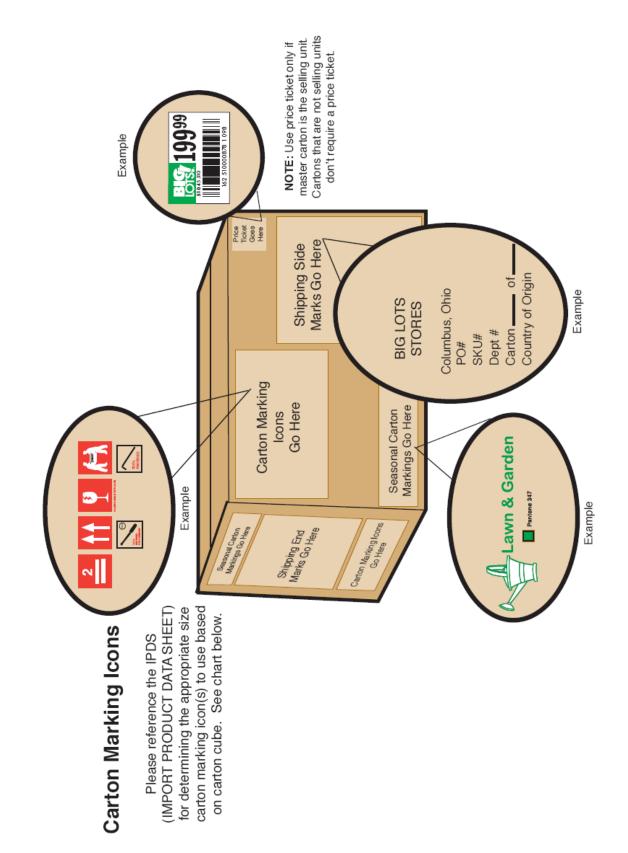
Figure 19: "Team lift" symbol

This symbol indicates that the weight of the container exceeds 70 lb or 31.75 kg and should <u>only</u> be lifted by 2 people.

Notes:

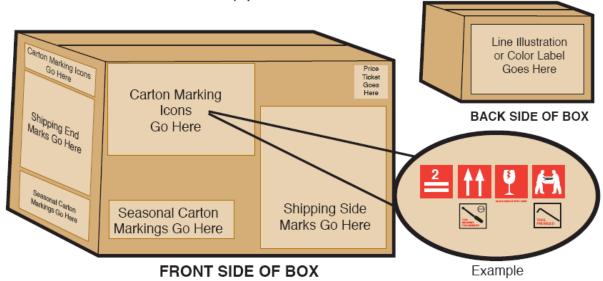
♦ This symbol should be located on at least 2 adjacent sides of the container and in Pantone 032.

	Vendor	DC	Store
Must be complied with by:	X	X	X



MASTER CARTON MARKING ICONS

TOOL(S) REQUIRED OR TOOL(S) PROVIDED ICONS



Carton marking icons are required to be placed on at least two different sides of the Master Shipping Carton. In the example above, the icons should be placed on the front, bottom right corner of the box when applicable. See <u>Carton Marking Icon Sizing Chart</u> for appropriate size carton marking icon(s).



TOOL REQUIRED FOR ASSEMBLY - Icon indicates what tool is required to assemble the item. Tells the customer/store associates what tool is required for assembly.



TOOL PROVIDED - Icon indicates what tool is included in box for assembly.

Some items may require more than one tool to assemble the item, see example below of how that should appear on box.

TOOLS REQUIRED FOR ASSEMBLY - Icon indicates what tools are required to assemble the item.

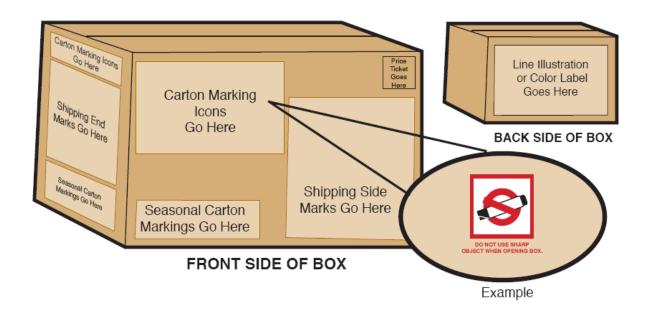






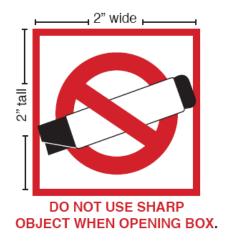


MASTER CARTON MARKINGS ICONS DON'T USE A KNIFE TO OPEN THIS BOX

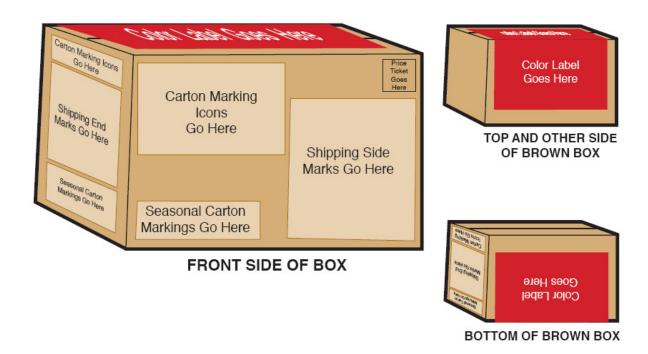


Carton marking icons are required to be placed on at least two different sides of the Master Shipping Carton. In the example above, the icon should be placed on the front, bottom right corner of the box when applicable. See <u>Carton Marking Icon Sizing Chart</u> for appropriate size carton marking icon(s).

This logo communicates to the stores that the merchandise inside (i.e. cushions, pillows, wicker furniture, etc.) can easily be cut and damaged when using a knife to open the box.



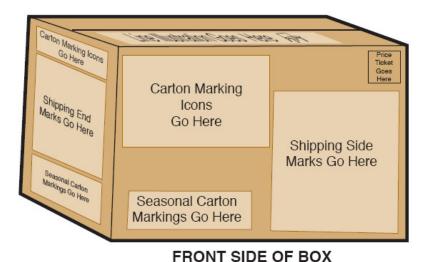
MASTER CARTON MARKINGS ICONS BROWN BOX WITH COLOR LABEL



This master shipping carton requires a color label to be placed on at least two different sides. In the example below, the shipping marks are placed on one end and one side panel of the box, as the front side has the color label.

IMPORTANT: The size of the color label placed on the front panel of a box must be at least 75% of the overall size of the front panel (front panel is considered the principal display panel). In the example above, a label has also been added to the side panel. Buyer may request this additional label if boxes are likely to be stacked in the store.

MASTER CARTON MARKINGS ICONS BROWN BOX WITH LINE ART





TOP AND OTHER SIDE OF BROWN BOX



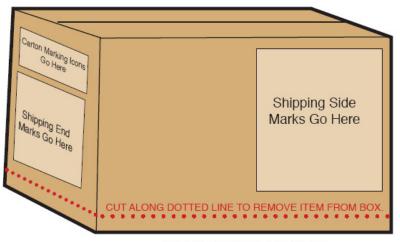
BOTTOM OF BROWN BOX

Carton marking line art is required to be placed on at least two different sides of the Master Shipping Carton. In the example above, the line art should be placed on the top and one side panel of the box when applicable.

This line art communicates to the customer/store associate what the merchandise looks like inside the brown box.

IMPORTANT: The size of the line art placed on the side panel of a box must be at least 75% of the overall size of the side panel. In the example above, the line art has also been added to the top panel. Buyer may request this additional label if boxes are likely to be stacked in the store.

MASTER CARTON MARKINGS FOR HEAVY AND/OR LARGE BULK BOXED ITEMS





BACK SIDE OF BOX

FRONT SIDE OF BOX

This carton marking is required to be placed on the bottom edge of the master shipping carton box when applicable. Please place dotted line 1" up from bottom of box.

For heavy and/or large bulk boxed items (i.e. curio cabinets, furniture, air compressors, generators, etc.), place dotted line across bottom of box, along with the following message:

CUT ALONG DOTTED LINE TO REMOVE ITEM FROM BOX

This will help customers/store associate to identify how to easily open the heavy box without damaging the item.

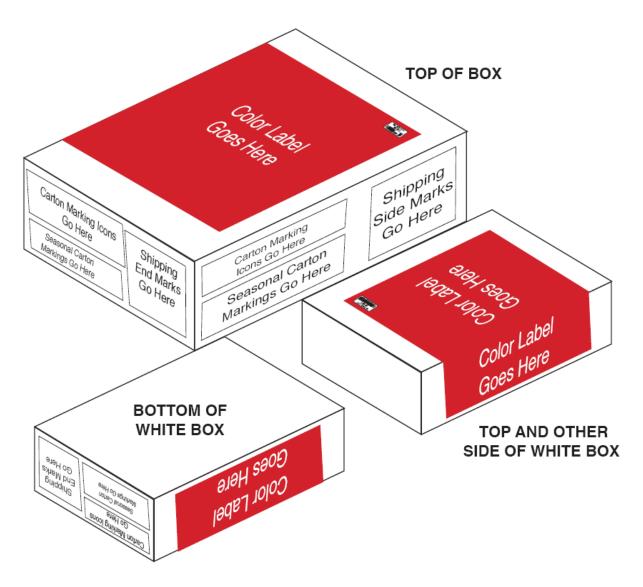
MASTER CARTON MARKINGS COLOR BOX

Some color boxes that are a case pack of one will also require carton markings to be placed on at least two different sides. In the example below, the shipping marks are placed on the bottom of the box and on the side panel of the color box. Graphics may need to be placed on the side panels to accommodate these carton markings.



MASTER CARTON MARKINGS WHITE BOX WITH WRAP-AROUND COLOR LABEL

This master shipping carton requires logos to be placed on at least two different sides. In the example below, the shipping marks are placed on one end and one side panel of the box, as the front side has the color label.



IMPORTANT: The size of the color label placed on the front panel of a box must be at least 75% of the overall size of the front panel (front panel is considered the principal display panel). In the example above, a label has also been added to the side panel. Buyer may request this additional label if boxes are likely to be stacked in the store.

MASTER CARTON MARKINGS WHITE BOX WITH WRAP-AROUND COLOR LABEL

In the example below the carton marking will not fit on the end of the box; therefore, we should place the carton marking on another dominant side panel, as the front side has the color label.

TOP AND OTHER SIDE OF WHITE BOX



TOP OF BOX



IMPORTANT: The size of the color label placed on the front panel of a box must be at least 75% of the overall size of the front panel (front panel is considered the principal display panel). In the example above, a label has also been added to the side panel. Buyer may request this additional label if boxes are likely to be stacked in the store.

Packaging Material Requirements - Minimum Standards

IMPORTANT! Big Lots vendors are responsible for conducting appropriate Primary Packaging performance testing to ensure that packaging does not fail in the supply chain or store. The material requirements provided below represent minimum standards.

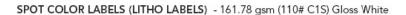
BACKER/BLISTER CARDS

- Lightweight items: 350 gsm (18 point), Sulfate Bleached Sheet (SBS) on both sides, C1S (Coated 1 Sided) on the Front with a UV Gloss Varnish.
- · Heavier items: Must be reinforced with a metal grommet at the hanging point.

COLOR BOXES

- Small items: 125 gsm (26#) E-Flute Singleface Kraft Corrugate + 250 gsm (12 point) Clay Coated News Back (CCNB) + UV Gloss Varnish.
- Larger items: 175 gsm (35#) E-Flute Singleface Kraft Corrugate + 350 gsm (26 point) Clay Coated News Back (CCNB) + UV Gloss Varnish.

NOTE: Packaging for very large and/or heavier items should be built to meet specific performance needs.



Corrugated Maximum Weight

- If box (L''+W''+D'') = up to 75" in size or 30 lbs in weight If box (L+W+D) = up to 190.5 cm in size or 13.6 kg in weight
- If box (L"+W"+D") = 75"-90" in size or 30 lbs 50 lbs in weight
 If box (L+W+D) = 190.5 cm 228.6 cm in size or 13.6 kg 22.6 kg in weight
- If box (L"+W"+D") = 90" 110" in size or 50 lbs 75 lbs in weight
 If box (L+W+D) = 228.6 cm 279.4 cm in size or 22.6 kg 34 kg in weight
- If box (L"+W"+D") = 110" 130" in size or 75 lbs 100 lbs in weight If box (L+W+D) = 279.4 cm 330.2 cm in size or 34 kg 45.3 kg in weight

Suggested Corrigated Box 32 ECT / 200# C-Flute (singlewall)

44 ECT / 275# C-Flute (singlewall)

48 ECT / 275# C/B-Flute (doublewall)

50 ECT / 350# C/B-Flute (doublewall)

• If box (L"+W"+D") = greater than 130" in size or greater than 200 lbs in weight 80 ECT / 350# C/B-Flute (doublewall with innerpack) If box (L"+W"+D") = greater than 330.2 cm in size or greater than 90.7 kg in weight Must meet ISTA Standards

DISPLAY BOXES (PDQ TRAYS)

- 32 ECT (Edge Crush) OR 200# (Burst/Mullen) B-Flute
- Minimum liner combination:

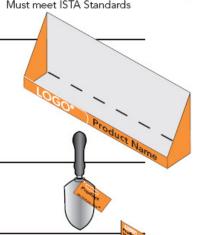
175 gsm (35#) liner x 112 gsm (23#) medium x 175 gsm (35#) liner

270 gsm (12 point) C2S (Coated 2 sided), Sulfate Bleached Sheet (SBS) on both sides, C2S (Coated 2 Sided) with a UV Gloss Varnish.

HEADER CARDS

HANG TAGS

- Lightweight items: 300 gsm (14 point) C2S (Coated 2 Sided) Sulfate Bleached Sheet (SBS)
- Heavier items: 350 gsm (18 point) or greater C2S (Coated 2 Sided) Sulfate Bleached Sheet (SBS)
- Extremely heavy items: Must be reinforced with a metal grommet at the hanging point.



10/07

Operating Procedures for Trade Community Regarding Implementation of the Wood Packaging Materials (WPM) Regulation

Background:

The USDA's APHIS has revised its import regulation for WPM, 7 CFR § 319. The final rule was posted in the Federal Register September 16, 2004, with an effective date of September 16, 2005.

The regulation requires regulated WPM used in international trade to be treated to kill harmful insects that may be present. WPM must be marked with the International Plant Protection Convention (IPPC) logo, the two-letter ISO code for the country that treated the WPM, the treatment facility number assigned by the national plant protection organization, and either the abbreviation HT (heat treatment) or MB (methyl bromide). The rule states that regulated wood packaging materials must be marked in a visible location on each article, preferably on at least two opposite sides of the article, with a legible and permanent mark that indicates that the article meets the new requirements. Paper treatment certificates will no longer be required or accepted. An example of an acceptable WPM mark is:



XX represents the ISO country code.

000 represents the unique number assigned by the national plant protection organization.

YY represents either HT for heat treatment or MB for methyl bromide fumigation.

The regulation restricts the importation of many types of wood articles, including wooden packaging materials such as pallets, crates, boxes, and pieces of wood used to support or brace product. The regulations currently refer to these types of wood packaging materials as solid wood packing materials, defined as "wood packing material other than loose wood packing material, used or for use with product to prevent damage, including, but not limited to, dunnage, crating, pallets, packing blocks, drums, cases, and skids." Effective September 16, 2005, the U.S. regulation allows non-compliant regulated WPM to be reexported. CBP recognizes that the usage of this term may be confusing. For purposes of CBP implementation of the USDA regulation, "reexport" will refer to the immediate export of violative WPM and, where the violative WPM cannot be separated from the accompanying product, the immediate export of the violative WPM and any accompanying product. By regulation, no treatment options for WPM being imported into the United States are available.

For the purposes of this rule, WPM imported as product, such as a container or truckload of new or unused pallets, will still be considered WPM and subject to the rule. Its status as product is irrelevant.

Non-regulated and Exempt Wood and Wood Products:

Regulated WPM do not include any manufactured items, such as worked wood items, even if those items are used to contain other non-regulated product. Examples of such non-regulated manufactured items might include such things as carved or formed wooden bottle stoppers, ammo crates, wooden boxes built to house fuel gauges or armaments, *etc*. Wine crates for any vintage year prior to 2006, are also non-regulated; wine crates for vintage year 2006 and beyond are regulated.

Regulated WPM do not include any manufactured wood, such as fiberboard, plywood, polywood, whisky and wine barrels, strand board, and veneers, nor do they include "loose wood packing materials" as defined in 7 CFR § 319.40-1. Examples of loose wood packing materials include excelsior (wood wool), sawdust, and wood shavings, produced as a result of sawing or

shaving wood into small, slender, and curved pieces. Dunnage is not always loose wood packing materials; when it is not, it is regulated.

The regulation allows importation without marking of otherwise-regulated WPM used by the U.S. Department of Defense ("DOD") to package non-regulated articles, including commercial shipments pursuant to a DOD contract.

By reciprocal regulations, WPM made from Canadian origin wood or U.S. origin wood (or a combination of Canadian origin wood and U.S. origin wood) will be exempt from treatment and marking under this regulation when used in trade between these two countries. For purposes of enforcement of this exception, and absent acceptable proof to the contrary, Customs will consider the country of origin of product coming from Canada to be the country of origin of the accompanying WPM.

The only remaining exemption for imports from Mexico permits importation of unmarked firewood, mesquite wood for cooking, and small, noncommercial packages of un-manufactured wood for personal cooking or personal medicinal purposes as long as these items arrive directly from Mexican Border States.

General Note 4(a) of the Harmonized Tariff Schedule of the United States (HTSUS)(2006) As of January 2006

The following countries, territories and associations of countries eligible for treatment as one country (pursuant to section 507(2) of the Trade Act of 1974 (19 U.S.C. 2467(2)) are designated beneficiary developing countries for the purposes of the Generalized System of Preferences, provided for in Title V of the Trade Act of 1974, as amended (19 U.S.C. 2461 et seq.):

Independent Countries

Afghanistan Croatia Lebanon Sao Tomé and Principe Albania Djibouti Lesotho Senegal Algeria Dominica Macedonia, Former Serbia and Montenegro Dominican Republic Yugoslav Republic of Angola Seychelles Antigua and Barbuda Ecuador Madagascar Sierra Leone Malawi Solomon Islands Argentina Egypt Armenia El Salvador Mali Somalia Bahrain **Equatorial Guinea** Mauritania South Africa Bangladesh Eritrea Mauritius Sri Lanka Barbados Ethiopia Moldova Suriname Belize Fiji Swaziland Mongolia Benin Gabon Mozambique Tanzania Bhutan Gambia, The Namibia Thailand Bolivia Nepal Togo Georgia Bosnia and Hercegovina Ghana Niger Tonga Botswana Grenada Nigeria Trinidad and Tobago **Brazil** Guatemala Oman Tunisia Pakistan Turkey Bulgaria Guinea Burkina Faso Guinea-Bissau Panama Tuvalu Burundi Guvana Papua New Guinea Uganda Haiti Cambodia Paraguay Uruguay Honduras Uzbekistan Cameroon Peru Cape Verde India **Philippines** Vanuatu Central African Republic Indonesia Romania Venezuela Chad Russia Republic of Iraq Yemen Colombia Jamaica Rwanda Comoros Jordan St. Kitts and Nevis Zambia Zimbabwe Congo (Brazzaville) Kazakhstan Saint Lucia Congo (Kinshasa) Kenya Saint Vincent and Costa Rica Kiribati the Grenadines Côte d'Ivoire Kyrgyzstan Samoa

GSP Indemnification Agreement

In consideration for its business dealings with Big Lots Stores, Inc. and/or its affiliated entities (collectively, "Big Lots"), the vendor listed below ("Vendor") agrees to the following:

Vendor hereby acknowledges receipt of the attached memorandum regarding the rules and regulations of the Generalized System of Preferences ("GSP") and certifies that the products which Big Lots purchases, from it qualify for duty free treatment under the GSP requirements of U.S. laws and regulations, and makes the following specific certifications:

- (1) The products are eligible products under the GSP;
- (2) The products are the product of a beneficiary developing country or an association treated as a single beneficiary developing country under GSP (in either case, a "BDC");
- (3) The products are to be shipped directly from the BDC to the United States, and
- (4) One of the following applies:
 - (a) The products are wholly the growth product or manufacture of a BDC. [Note: If the products are wholly the growth, product or manufacture of a BDC, a statement to that effect shall be included on the commercial invoice.]
 - (b) At least 35% of the appraised value of the products is attributable to (i) the cost or value of the materials produced in a BDC and (ii) the direct costs of processing in a BDC. [Note: If the products are not wholly the growth, product or manufacture of a BDC, the Vendor shall submit supporting documents within twenty (20) days of a request from Big Lots or U.S. Customs.]
- (5) Vendor will be compliance with all GSP requirements and obligations with respect to the products.
- (6) Vendor will maintain supporting production, manufacturing and financial documents for a period of five (5) years from the date of exportation. Vendor shall submit supporting documents within twenty (20) days of a request from U.S. Customs or Big Lots. Supporting documents are inclusive of, but not limited to:
 - (a) The manufacturer-completed GSP worksheet (the form of which is attached hereto) detailing the allocation of costs, the general ledger accounts and other documentation that supports the GSP claim.
 - (b) Bills of materials, trial balances, general ledgers, raw material invoices, purchase history reports, inventory records, freight bills, insurance premiums, packing invoices, sales invoices relating to waste shipments, tax bills, broker invoices, cost sheets, rent bills, utility bills, financial statements and any other documentation which may be required by U.S. Customs to validate the GSP claim.
- (7) In addition to the obligations set forth elsewhere in this Agreement, the Vendor will otherwise cooperate fully with Big Lots in providing appropriate information, documentation, and certification in response to any provider inquiry or audit (i.e., Customs, a court of law, or other binding authority).
- (8) Vendor will complete a GSP Declaration in the format annexed for each qualifying shipment and, at the time of shipment, provide an original to Big Lots together with the commercial invoice, packing list, bills of lading and other documentation upon which Big Lots' payment is conditioned.

In the event that it is determined by Customs, a court of law, or other binding authority that the products shipped and entered based upon a GSP Declaration executed by the Vendor do not qualify for GSP, Vendor shall indemnify and hold harmless Big Lots Stores, Inc., its parent and its affiliates and the directors, officer and agents of each for any duties and penalties that are assessed on such products, as well as all other liabilities damages, costs and expenses (including legal fees) incurred in connection with or related to such matter, audit, or proceeding.

By signing below, you represent that you are a representative of the Vendor, you have (on the Vendor's behalf) read and understand this GSP Indemnification, and are duly authorized to strictly bind the Vendor to the terms of this GSP Indemnification Agreement. Please fax or scan this page with your signature to our Global Sourcing Specialist at globalsourcing@biglots.com.

Vendor's Full Name	
Signature of Vendor's Authorized Representative	
Printed Name of Vendor's Authorized Representative	
Title of Vendor's Authorized Representative	
Date	
Fax:	

G	SP DECLARATION
EMAIL: FAX:	
ARTICLE NAME, DESCRIPTION AND ITEM NO.:COUNTRY OF ORIGIN:	
MAJOR CONSTITUENT MATERIALS MATERIAL 1. 2.	COUNTRY OF ORIGIN OF MATERIAL
3 4. 5.	
BRIEF DESCRIPTION OF PROCESSING OPERATION	NS PERFORMED BY VENDOR:
CONFIRMATION OF GSP ELIGIBILITY BY	, ON BEHALF OF IALS USED TO PRODUCE THE ARTICLE ORIGINATE IN THE
	NT OF THE SALES PRICE OF THE ARTICLE CONSISTS OF THE SUM COUNTRY OF ORIGIN AND DIRECT COSTS OF PROCESSING
3. VENDOR AGREES TO MAINTAIN ALL BUSINES QUALIFY FOR THE GSP FOR A PERIOD OF 5 YEAR	S RECORDS NECESSARY TO ESTABLISH THAT THESE ARTICLES AS FROM DATE OF EXPORT.
	ECORDS NECESSARY TO ESTABLISH GSP ELIGIBILITY TO BIG IS OR U.S. CUSTOMS WITHIN 20 DAYS OF REQUEST.
BIG LOTS IS REQUIRED TO PAY DUTIES OR PENA	ACCORDANCE WITH THE GSP INDEMNIFICATION AGREEMENT IF LITES OR INCUR OTHER EXPENSES BY REASON OF VENDOR'S TE DOCUMENTATION CONFIRMING GSP ELIGIBILITY IN A
I DECLARE THAT THE INFORMATION PROVIDED	IN THIS DOCUMENT IS TRUE AND CORRECT:
SIGNATURE:N	AME AND TITLE:
DATE:	
This form must be completed and submitted with the fina	l IPDS to the Global Sourcing Specialist.

Page 1	Source Record of Cost Example of Potential Records to Support Cost GSP Declaration, Bill of Materials (BOM), Trial	Balance, G/L, Vendor Invoices, Purchase History Reports, Inventory Records	Records BOM, Trial Balance, G/L	Freight Bills, BOM, Trial Balance, G/L	Insurance Premiums, BOM, Trial Balance, G/L	Packing Invoices, BOM, Trial Balance, G/L	Sales invoices relating to waste shipments BOM, Trial Balance, G/L, Tax Bills, Broker	Invoices		GSP Declaration, Cost Sheets, BOM, G/L	GSP Declaration, Cost Sheets, BOM, G/L	GSP Declaration, Cost Sheets, BOM, G/L	GSP Declaration, Cost Sheets, BOM, G/L	G/L, Trial Balance, Rent Bills	G/L, Trial Balance, Utility Bills
	General So Ledger Re Chart of of Account (Re		#######################################	#####	#####	#####	#####	#####		######	#####	## ## #	######	#####	#####
Article # / SKU	Description of Cost		Vendor's actual cost of materials Purchase price or material variances	Cost Incurred in transporting material to the vendor	Cost incurred in transporting material to the vendor	Cost incurred in transporting material to the vendor	Net cost of any waste or spoilage Duties and/or taxes imposed on the materials by	the BDC	Employee costs directly involved in production of finished good on the job training transportation	cost, insurance compensation, Fringe Payroll Benefits, Housing Allowance, Severance, Social	Security Taxes, Medial Insurance Expenses, Bus Transportation	roust or employees who receive, unload, and socon raw materials in manufacturer's plant, distribute materials to assembly line, maintain storage area and raw material inventory records, and pack and	prepare articles for shipment Compensation, including fringe benefits, to the extent these personnel functions are directly involved in the production of the specific	Rent attributable to that portion of the building space directly used in the processing operations	Cost of utilizes, such as electricity, telephone, and water to the extent they are actually used in the production process of the merchandise
	Value of Non-Direct Processing Cost	abor, Direct						-	_						
HEET	Value of Direct Processing Costs	als, Direct La	\$\$\$\$\$ \$\$\$\$\$\$	\$\$\$\$\$	\$\$\$\$\$	\$\$\$\$\$	\$\$\$\$\$\$	\$\$\$\$\$		\$\$\$\$\$	\$\$\$\$\$\$	\$\$\$\$\$	\$\$\$\$\$\$\$	\$\$\$\$\$	\$\$\$\$\$
GSP WORKSHEET	Direct Processing Operation Cost	Product Cost (Materials, Direct Labor, Direct Overhead)	Materials Materials Variances	Freight	Insurance	Packing	Scrap (Net)	Duties & Taxes	Direct Labor	Direct labor	Quality Control	Shipping & Receiving	Other Employees (Supervisory, Engineering, Plant	Direct Overhead Rent	Utilities

GSP WORKSHEET		Article # / SKU		Page 2
Period Cost		Cost of renting, repairing, maintaining, modifying		
Equipment \$\$\$\$\$		production machinery	####	G/L, Trial Balance
Indirect Materials \$\$\$\$\$		Cost of materials consumed in production such as		
		lubricants, chemicals fuel	####	G/L, Trial Balance
		Dies, molds, tooling (this category can also include		
Fixed Assets \$\$\$\$\$		Assists) whether capitalized or expensed	#####	G/L, Trial Balance
		Depreciation on machinery and equipment which		
Depreciation \$\$\$\$\$		are allocable to the merchandise	####	G/L, Trial Balance
Total Period Cost \$\$\$\$\$				
Other Cost		Packing performed in BDC and essential for the		
Packaging \$\$\$\$		shipment of an eligible article to the US	#####	G/L, Trial Balance, Invoices for Packaging
		Pro-rate share of taxes on the part of the building		
Taxes		used in the processing operations	####	G/L, Trial Balance, Tax Bills
		Cost of property insurance covering machinery		
Insurance		and equiment used in the production process	#####	G/L, Trial Balance, Insurance Premiums
Total Other Costs 88888				
Non-Direct Processing Cost				
Indirect Materials	88888	Office supplies, stamps	#####	G/L, Trial Balance
Indirect Labor	\$555	Labor costs incurred for indirect labor such as administrative salaries, sales people, accounting	#####	G/I Trial Balance
		personnel		
		Rent and Utilizes on the portion of the building		
General Expenses	\$\$\$\$\$	used for personnel offices, accounting department,	#####	G/L, Trial Balance, Rent & Utilities
		and other administrative functions		
Taxes	88888	Sales Tax	#####	G/L, Trial Balance, Tax Bills
Insurance	88888	Casual and liability Insurance	#####	G/L, Trial Balance, Insurance Premiums
Profit	\$\$\$\$\$	Vendor Profit/Loss	####	Financial Statements
		的 1000 1000 1000 1000 1000 1000 1000 10		
Total Operation Cost \$\$\$\$\$	\$\$\$\$\$		10 10	

Total Value of Direct Processing Cost

35%

Value Declared to customs (Actual Cost)

(Section 6.2- Ross Stores method of valuation is Transaction Value, which is the price actually paid for the imported merchandise to the vendor)
*Period Costs- The allocation of these cost must be pro-rated against production records for specific items or over specific product runs over specific SKUs

GSP MEMORANDUM

THIS MEMORANDUM IS NOT INTENDED TO PROVIDE AND SHALL NOT BE CONSTRUED AS PROVIDING LEGAL ADVICE. YOU SHULD CONSULT YOUR OWN ATTORNEY FOR ADVICE.

A. INTRODUCTION

This memorandum provides an overview of the requirements associated with the Generalized System of Preferences ("GSP") program administered by the U.S. Customs Service. Under the GSP program, selected articles from designated beneficiary developing countries ("BDC's") may qualify for preferential (generally duty-free) treatment. The list of articles and countries eligible under the program is periodically reviewed by the United States and is subject to amendment. The list of BDC's appears in General Note 4(a), Harmonized Tariff Schedule of the United States ("HTSUS"). However, whether an article from a BDC is GSP-eligible depends on the tariff classification of that article under the HTSUS. Therefore, the correct tariff classification provision must be ascertained, and the tariff provision will indicate whether such article is GSP – eligible. The "Special" rate of duty column associated with the tariff provision in the HTSUS will indicate the symbol "A" if the product is GSP-eligible. If the A is followed by an asterisk ("A*"), the article is GSP-eligible *except* from certain BDC's. Those excepted BDC's are listed in General Note 4(d), HTSUS.)

In order for an eligible article from a designated beneficiary developing country (hereinafter, "BDC") to qualify for GSP treatment, three principal legal requirements must each be satisfied:

- 1. The article must be considered a "product of" the BDC;
- 2. The sum of (1) the cost or value of the materials produced in the BDC or any 2 or more countries which are members of the same association of countries which are treated as one country under the GSP program, plus (2) the direct cost of processing operations performed in such BDC or such member countries, is not less than 35 percent of the appraised value of such article at the time of its entry into the customs territory of the United States; and
- 3. The article is imported into the United States directly from the BDC.

See General Note 4(c), HTSUS. Each of these legal requirements is discussed in further detail below. Reference will also be made to the Customs Regulations, which appear in 19 C.F.R. 10.171 - 10.178.

B. GSP-ELIGIBILITY REQUIREMENTS

1. "Product of" Requirement

An article either must be wholly the growth, product or manufacture of the BDC or manufactured in an operation resulting in the finished article having the BDC as its legal country of origin for U.S. Customs purposes. In other words, the article must be a "product of" the BDC for U.S. Customs purposes. See 19 C.F.R. 10.176.

When an article is produced from non-BDC materials, the production in the BDC must normally qualify as a "substantial transformation" in order for the finished article to be considered a product of the BDC. Determinations as to whether a substantial transformation has occurred are made on a case-by-case basis depending upon the nature of the product and the complexity of the manufacturing operations.

In general, a substantial transformation occurs if (1) a new and different article of commerce emerges (2) as the result of a substantial manufacturing or processing operation. Customs has stated that a new and different article of commerce is an article that has undergone a change in: commercial designation or identity, fundamental character, or commercial use. The factors to be considered in determining whether the article has been subjected to a substantial manufacturing or processing operation include: the physical change in the article as a result of the processing, the time involved in processing, the complexity of the processing, the level of degree or skill and/or technology required, and the value added to the article.²

¹ Certain articles cannot be designated as GSP-eligible. See General Note 4(c), HTSUS. Non-eligible articles include: textile and apparel articles which are subject to textile agreements; certain watches; import-sensitive electronic articles; import-sensitive steel articles; generally, footwear, handbags, luggage, flat product, work gloves, and leather wearing apparel.

The fact that an article satisfies the 35 percent value-added requirement discussed in the next section, however, does not automatically mean that the article is a "product of" the BDC.

2. 35 Percent Value Content Requirement

In order for an otherwise eligible article to qualify under GSP, the sum of (1) the cost or value of the materials produced in the BDC or any 2 or more countries which are members of the same association of countries which are treated as one country under the GSP program, plus (2) the direct cost of processing operations performed in such BDC or such member countries, is not less than 35 percent of the appraised value of such article at the time of its entry into the customs territory of the United States.³

a. Materials

The concept of "materials produced" in the BDC includes not only materials which are wholly the growth, product or manufacture of the GSP country, but also foreign materials imported into the BDC that undergo a "double substantial transformation" in the BDC (*i.e.*, one substantial transformation into a new and different "intermediate" article of commerce that is then subjected to a second substantial transformation into a new and different finished article). As discussed above, determinations as to whether a substantial transformation has occurred are made on a case-by-case basis depending upon the nature of the product and the complexity of the manufacturing operations.

However, in order to effectuate the GSP program, U.S. Customs may permit imported materials to count toward the 35 percent value-added requirement by liberalizing its substantial transformation test. For instance, if raw material is imported into the BDC and substantially transformed into components or parts, which are then assembled to form the final product, U.S. Customs may permit the final assembly to qualify as the second (double) transformation even though by itself the assembly might not qualify as a substantial transformation. In such case, the imported materials must be transformed into "intermediate" articles of commerce, and the subsequent assembly must be meaningful and result in yet a new and different final article of commerce. See Customs Ruling HQ 557284 (1993), citing *Texas Instruments. Inc. v. United States*, 681 F.2d 778 (Fed. Cir. 1982).

In order to calculate the cost or value of qualifying BDC materials, the Customs Regulations at 19 C.F.R. 10.177(c) provide the following guidelines:

- (c) Determination of cost or value of materials produced in the beneficiary developing country.
- (1) The cost of value of materials produced in the beneficiary developing country includes:
 - (i) The manufacturer's actual cost for the materials;
 - (ii) When not included in the manufacturer's actual cost for the materials, the freight, insurance, packing, and all other costs incurred in transporting the materials to the manufacturer's plant;
 - (iii) The actual cost of waste or spoilage (material list), less the value of recoverable scrap; and
 - (iv) Taxes and/or duties imposed on the materials by the beneficiary developing country, or an association of countries treated as one country, provided they are not remitted upon exportation.⁴
 - (2) Where the material is provided to the manufacturer without charge, or at less than fair market value, its cost or value shall be determined by computing the sum of:
 - (i) All expenses incurred in the growth, production, manufacture or assembly of the material, including general expenses;

³ General Note 4(a) contains a list of various associations of countries treated as one country. For example, ASEAN consists of Cambodia, Indonesia, Philippines, and Thailand. Another example is the Andean Group, which consists of Bolivia, Colombia, Ecuador, Peru, and Venezuela.

⁴ The cost of transporting non-qualifying materials to the manufacturer's plant does <u>not</u> count toward the 35 percent requirement.

- (ii) An amount for profit; and
- (iii) Freight, insurance, packing, and all other costs incurred in transporting the materials to the manufacturer's plant.

If the pertinent information needed to compute the cost or value of the materials is not available, the appraising officer may ascertain or estimate the value thereof using all reasonable ways and means at his disposal.

b. <u>Direct Costs of Processing Operations</u>

The Customs regulations (at 19 C.F.R. 10.178) provide that the "direct costs of processing operations" means those costs either directly incurred in, or which can be reasonably allocated to, the growth, production, manufacture, or assembly of the specific article under consideration. Such costs include, but are not limited to:

- (1) All actual labor costs involved in the growth, production, manufacture, or assembly of the specific article, including fringe benefits, on-the-job training, and the cost of engineering, supervisory, quality control, and similar personnel;
- (2) Dies, molds, tooling, and depreciation on machinery and equipment which are allocable to the specific product;
- (3) Research, development, design, engineering, and blueprint costs insofar as they are allocable to the specific product; and
- (4) Costs of inspecting and testing the specific product.

Items which are not included in the "direct costs of processing operations" are those which are not directly attributable to the product being imported. These include profit and general expenses of doing business which are either not allocable to the specific product or are not related to the growth, production, manufacture, or assembly of the product, such as administrative salaries, casualty and disability insurance, advertising, and salesman salaries, commissions or expenses.

On the other hand, all costs directly associated with the processing of the product, such as direct overhead, may qualify. In addition to the Customs regulations cited above, there have been various rulings issued by Customs which explain the differences between qualifying and non-qualifying labor costs and overhead for the purposes of satisfying the 35 percent value-added requirement. These differences are explained below.

(1) **QUALIFYING LABOR COSTS**

Wages and other costs of production personnel, quality control personnel, engineering personnel, and supervisory production personnel.

Whether a particular labor cost constitutes a direct cost of processing is generally determined by the function the employee performs. HQ 555043 (1988). Where costs are incurred for both qualifying and non-qualifying functions, the costs must be allocated among production and non-production functions. This allocation must be done on per model basis. The allocation must also have a rational basis and conform to Generally Accepted Accounting Principles.

Set forth below are some examples of specific labor costs, beyond those directly associated with assembly line workers, that can qualify toward the 35% local-value content requirement.

Generally, all compensation to production workers including first-line supervisors and quality control inspectors, directly involved in the production of the product may be included in the 35% value-content as a direct cost processing. C.S.D. 80-246 (1980); HQ 541689 (1978).

Compensation of group leaders, quality control supervisors and manufacturing foreman are direct costs of processing to the extent that they perform as first line supervisors of workers directly involved in the production process. HQ 554246 (1987).

Additionally, compensation to materials handlers, shipping and receiving personnel, packers and similar personal also qualify to the extent they handle the specific parts for the model. C.S.D. 80-208 (1980) (cost of employees who receive, unload, and stock raw materials in manufacturing

plant, distribute materials to assembly line, maintain storage area and raw material inventory records, pack and prepare eligible articles for shipment also constitute a direct cost of processing).

Export packaging costs for the final product, including the cost of packaging materials if BDC-origin, are included. C.S.D. 79-199; HQ 047532 (1977).

The transportation costs, if any, of transportation provided to <u>direct labor employees</u> is a direct cost of processing. HQ 544067 (1989); HQ 554246 (1987) (Expenses incurred in transporting personnel to and from the factory to render services are directly related to the production process and are included as a direct cost of processing).

Compensation of janitorial personnel to the extent incurred in the production plant or factory area may be included. C.S.D. 80-208 (1980)

Compensation to maintenance personnel to the extent they maintain equipment used in the production of the product and not claimed under another heading may be included. HQ 543748 (1987).

Compensation of engineering personnel to the extent they provide services facilitating the production process may be included. HQ 544067 (1989). Where engineering services are provided to the manufacturer, the value of that assist may be included in the direct cost of processing. C.S.D. 80-208 (1980). Additionally, the cost of travel necessary to bring the engineer to the facility to render the services is similarly included.

The contract amount paid to and unrelated labor service firm that obtains the required labor for the production facility was found to be a qualifying labor cost. HQ 543748 (1987).

Fringe benefits

To the extent that qualifying labor costs do not include the cost of fringe benefits, those costs may separately qualify. Christmas bonuses, vacation pay, social security taxes, group insurance medical benefits, cost of a nursery, housing allowances, payroll taxes, and cost of a cafeteria or daily bus transportation provided to production personnel are qualifying costs. C.S.D. 80-246 (1980); HQ 555198 (1989); HQ 555379 (1989); HQ 555316 (1989); HQ 555043 (1988); HQ 543748 (1987); HQ 541689 (1978); HQ 541215 (1977).

Training Costs

Additionally, all on-the-job training costs for production personnel are also qualifying costs. 19 C.F.R. 10.178(a)(1).

(2) NON-QUALIFYING LABOR COSTS

Wages and other costs of non-production personnel, sales, managerial, accounting, etc.

Again, the function of the employee will generally determine if costs associated with that employee are non-qualifying costs. Generally, costs of personnel not directly associated with production are non-qualifying costs. Set forth below are some examples.

Costs associated with employees who perform <u>only</u> administrative functions, such as the general manager, personnel manager, plant security or accounting or payroll personnel are not considered direct cost of processing. HQ 544067 (1989); HQ 543748 (1987); C.S.D. 80-208 (1980); HQ 541249 (1977).

Compensation of the plant manager is not considered a direct cost of processing because he performs primarily administrative functions. However, Customs has held that to the extent the

plant manager performs first-line supervisor functions, the cost of his compensation may be claimed. HQ 543748 (1989). Thus, for a few employees, it may be possible to allocate compensation between qualifying and non-qualifying costs.

Fringe Benefits; Training Costs

As indicated above, all compensation, including fringe benefits and training costs of employees whose compensation is non-qualifying is also non-qualifying. This includes the cost of their automobiles and travel. HQ 543748 (1987); HQ 541249 (1977); C.S.D. 80-208 (1980); HQ 541215 (1977).

(3) **QUALIFYING DIRECT OVERHEAD**

Generally, all costs directly associated with the processing of the subject product can be claimed as qualifying direct overhead. Customs has indicated that without sufficient detail Customs will be unable to accept the overhead costs as qualifying costs. Where costs are incurred for both qualifying and non-qualifying functions, the costs must be allocated among production and non-production functions. This allocation must be done on per model basis. Again, the allocation must have a rational basis and conform to Generally Accepted Accounting Principles.

Design, engineering, R&D costs

Customs has held that where these items are provided to the manufacturer free of charge, they constitute an assist. The value of this assist must be declared on importation if not included in the price paid or payable. However, this assist also qualifies as a direct cost of processing. I.A. 83/76, HQ 541249 (1977); I.A. 41/76, HQ 541215 (1977); HQ 541080. The country of origin of such an assist is irrelevant. Caveat: To the extent engineering wages and similar personnel compensation are claimed as a labor cost, they should not be also claimed as an engineering expense.

The cost of an engineering service which provided installation and repair of equipment and parts, monitored, inspected and tested equipment and performed similar functions was found to be a qualifying cost <u>except</u> for that portion of the contract that represented the performance of administrative functions. HQ 543748 (1987).

Equipment rental fees

The equipment must be used in production in order for the fees to qualify. C.S.D. 79-63.

Equipment Depreciation

Depreciation on equipment and machinery directly involved in the production process are qualifying costs. HQ 555379 (1989); C.S.D. 80-246 (1980); I.A. 41/76, HQ 541215 (1977).

Repair and maintenance cost

The cost of maintenance, modification and repair of machinery used in the production process is a qualifying cost. C.S.D. 80-246 (1980); HQ 541215 (1977). The cost of the chief mechanic and other mechanics are qualifying costs to the extent they are maintaining the equipment. HQ 555316 (1989). However, to the extent the cost of these personnel are claimed as a qualifying labor cost, their cost must not be counted twice.

Molds, dies or tooling depreciation

Depreciation on tools, dies and molds directly involved in the production process are qualifying costs. I.A. 41/76, HQ 541215 (1977). Again, however, if not included in the invoice price, the value of the molds, dies, and tools are also assists if provided free of charge to the manufacturer, and must be added to dutiable value. If included in dutiable value, the value of these assists will qualify as a direct cost of processing regardless of the country of origin of the assist. HQ 541249 (1977); HQ 541080.

Inspection and Testing (if not previously included as a qualifying labor cost above)

Rent (apportioned between manufacturing and non-manufacturing space)

Rent on that portion of the building space directly used in the processing operations is considered a qualifying cost. C.S.D. 79-63; HQ 543748 (1987); I.A. 83/76, HQ 541249 (1977); HQ 555379 (1989).

Utilities (electricity and water, etc. apportioned to manufacturing)

The cost of fuel, water and electricity are direct costs of processing to the extent they are used in the production process. C.S.D. 79-63; C.S.D. 80-208 (1980); C.S.D. 80-246 (1980). For example, electricity used to operate machinery in the production process and the electricity used to light the work area are direct costs of processing.

Local Taxes (apportioned to manufacturing)

Taxes on that portion of the building used in the production process are a qualifying cost. C.S.D. 79-63; C.S.D. 80-208 (1980); HQ 056936 (1978); I.A. 41/76, HQ 541215 (1977).

Royalty fees

Certain royalty fees qualify. Customs has ruled that where engineering, design and production information is made available from the parent for the production of the product, royalties paid on a per-unit basis and reflected on the manufacturer's books as an operating expense constitute a "direct cost of processing". Headquarters Ruling Letter 543155 (1983). Similarly, where the royalty payments are made for on-site quality audits and/or for research and design of the specific model they qualify. C.S.D. 79-312 (1979).

Miscellaneous expenses directly related to processing the subject model

Certain other expenses may be claimed as a direct cost of processing. A review of the exact expenses involved is necessary to reveal whether any of these expenses may be claimed as qualifying. Set forth below are some examples.

Cost of property insurance covering machinery and equipment used in the production process are qualifying costs. HQ 543748 (1987); HQ 555379 (1989).

Local licensing fees required to manufacture the subject product are a qualifying overhead cost. I.A. 134/77, HQ 541689 (1978).

Telecommunication costs incurred to facilitate the inspection of product and first line supervision of the production process is a qualifying cost. HQ 555379 (1989); HQ 554246 (1987).

The interest expense on the acquisition of plant equipment and machinery is a direct cost of processing because it is a necessary expense directly related to the cost of machinery. C.S.D. 79-312 (1979).

(4) NON-QUALIFYING OVERHEAD

Selling Expenses (travel, etc.)

Office Expenses (telephone, stationary, office equipment, etc.)

General office expenses such as postage or telecommunication charges are not qualifying costs. 19 C.F.R. 10. 178(b)(2); HQ 554067 (1989); HQ 543748 (1987); HQ 541215 (1987), HQ 055694 (1979).

Advertising

Entertainment

Utilities (electricity and water apportioned to non-manufacturing)

The cost of fuel, water and electricity are not direct costs of processing to the extent they are not used in the production process but support administrative functions. C.S.D. 80-246 (1980). For example, electricity used for lighting and air conditioning the administrative office would not be qualifying costs. Again, however, those same costs would qualify if incurred with respect to that portion of the plant dedicated to production.

Rent (apportioned to non-manufacturing)

Rent on that portion of the building space used for the accounting department, personnel offices and other administrative functions are not considered a qualifying cost. HQ 541249 (1987); HQ 555379 (1989).

Fire and Casualty Insurance

Fire, casualty and liability insurance are not direct costs of processing. 19 C.F.R. 10.178(b)(2); HQ 555316 (1989); HQ 555379 (1989).

Miscellaneous expenses not directly related to processing the subject model

Again, a careful review of the so-called "other expenses" is necessary. Set forth below are some examples of miscellaneous costs that do not qualify.

The cost of automobiles is not a direct cost of processing. HQ 555379 (1989).

On-site medical personnel costs are not a qualifying cost even if required by law for production personnel. C.S.D. 80-208 (1980).

Legal expenses. C.S.D. 80-208 (1980).

3. "Direct Importation" Requirement

Eligible articles must be "imported into the customs territory of the United States directly from" the BDC. While certain exceptions apply, a satisfaction of the "imported directly" criterion normally requires direct shipment from the BDC to the United States without passing through the territory of any other country. If the shipment is through the territory of any other country, the product may not enter into the commerce thereof while en route and the invoice, bills of lading, and other shipping documents must normally show the U.S. as the final destination. Shipment of a product from a BDC to an inventory warehouse in a third country would likely not satisfy the direct importation requirement.

Special attention must be paid to instances in which a product of a BDC is further processed in a foreign country. In such instances, Customs will require that it be returned to the BDC for further processing so as to reenter the commerce of the BDC prior to shipment to the United States. In other words, the return to the BDC must consist of more than a mere pass-through. At the very least, the returned articles must be inspected and packaged in the BDC.

C. CONCLUSION

As discussed, the 35 percent value content requirement may be satisfied through eligible materials alone, through direct processing operations alone or through a combination of the two. In conclusion, we will discuss below two examples to illustrate this principle.

Example 1 – Costs of Direct Processing Alone

Facts: Foreign materials are shipped to a BDC where they are manufactured into a product of the BDC, but do not undergo a "double substantial transformation". The actual labor cost involved in the production of the finished article in the BDC is 40% of the total U.S. Customs appraised value of the finished product.

Conclusion: The qualifying GSP costs are 40%. The 35% value content requirement is satisfied through the labor costs alone.

Example 2 - Costs of Materials Plus Direct Processing

Facts: Foreign materials are shipped to a BDC where they are manufactured into a product of the BDC. The foreign materials, which represent 20% of the total U.S. Customs appraised value of the finished product, undergo a "double substantial transformation" in the BDC. The actual labor costs involved in the production of the finished article in the BDC is 25% of the total U.S. Customs appraised value

Conclusion: The qualifying GSP costs are 45%. The 35% value content requirement is satisfied through the materials cost (which may be included in this example because there is a double substantial transformation) plus the labor cost.

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NEW DELHI

Note: All New Delhi cargo should be delivered to Mumbai for shipment and/or consolidation

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**Booking for Shenzhen and Huangpu should be sent to NYKL Hong Kong directly, where assistance will be provided in arranging shipments

CONTAINER LOADING PLAN - BIG LOTS STORES, INC. AND ITS AFFILIATES

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To: NYK Logistics FM: (Shipper):	Shipper Address:	CONTAINER NO.	PORT OF LOADING	SHIPPING MARKS									

1. Please submit one Container Loading Plan per container.

2. HTS# is the Harmonized Tariff Schedule (same as Tariff# listed in merchandise information sheet).

3. Please submit the Container Loading Plan with accurate details before the Container Loading Plan cutoff time. Any missing or incomplete information will result in either cargo detention at the loading port or be rejected by U.S. Customs.

							(5) ETA Yantian	
(1) Carrier's CY Accepting Date		(2) NYK CLP Curoff Date		(3) Carrier's SI Cutoff Date		(4) Carrier's CY Cutoff Date		(6) ETD Yantian
φ		φ	-5	4	ώ	-2	-	0
18-Jan	19-Jan	20-Jan	21-Jan	22-Jan	23-Jan	24-Jan	25-Jan	26-Jan
Sunday	Monday	Tuesday	Wednesday	Thursday	Friday	Saturday	Sunday	Monday
		7	7	0				
(1) Carrier's CY Accepting Date	iting Date		18-Jan	Container starts accepting Laden Containers for export to USA at Yantian CY	pting Laden Cont	tainers for export to U	SA at Yantian CY	
(2) NYK CLP Cutoff Date 20-Jan	ate 20-Jan		20-Jan	Vendor must submit CLP (Container Load Plan) to NYK	CLP (Container L	Load Plan) to NYK		
(3) Carrier's SI Cutoff Date	Date		22-Jan	NYK must submit SI (Shipping Instruction) to Carrier	Shipping Instruc	tion) to Carrier		
(4) Carrier's CY Cutoff Date	Date		24-Jan	Vendor must delivery	Laden Containe	Vendor must delivery Laden Container to carrier's CY for export to USA	rport to USA	
(5) ETA Yantian			25-Jan	Container vessel arrives the Yantian port	es the Yantian p	iort		
(5) ETD Yantian			26-Jan	Container vessel departures Yantian for export to USA	artures Yantian fi	or export to USA		
Three Citoff Dates								
(2) NYK CLP Cutoff Date	ate		NYK CLP Cutoff	NYK CLP Cutoff Date is 2 days prior to Carrier's SI Cutoff Date	Carrier's SI Cut.	off Date		
(3) Carrier's SI Cutoff Date	Date		Carrier's SI Cutol	Carrier's SI Cutoff Date is normally 3 days prior to Vessel ETA (Estimated Time Arrival)	ys prior to Vesse	el ETA (Estimated Tin	ne Arrival)	
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Document Delay Notice



** DOCUMENT DELAY NOTICE ** <u>BIG LOTS STORES, INC.</u> (Pakistan Shipment)

	(Pakist	an Shipment)			
			I	Ref. no.	
Messers:		21	st DDN : nd DDN	Send Send Send	
Dear Sir/Madam;					
Further to our telephone condelay of document. You are with all shipping documents of delay, please describe on this p	requested to submit NYK of the following shipments a	L (ORIGIN), within 2 already effected 2 days	4 hours	after this notice i	s served,
Vessel/Voyage	Sailing Date	P.O. # / Stock #		Pkgs	Reasons
In the event you fail to supply any reasonable reason of delay	•	•			vise us of
Days following sailing: Fee: US	6-7 days 8-9 \$\$50/set equivalent US\$		0 days + JS\$63/se	t equivalent	
These fees are charged for the expenses for sending delay rathention.					
Very truly yours,					



NYK Logistics

e-Booking Registration Form

Please complete this form and forward to nlhk_biglots@hk.nyklogistics.com

If available, please provide Big Lots PO number _	for our internal reference
Consignee	Internal Use Only
Big Lots Stores, Inc.	Vendor Code:
Company Information - To be declared as the sh	ipper of booking.
Company Name: Address 1: Address 2: City: Country:	
Administrative User Information – Manages org passwords information.	ganization's User profile and
Login ID: Full Name: Phone Number: Email Address: Fax Number:	
Additional User Information – Responsible for p	erforming eBooking operation.
Login ID: Full Name: Phone Number: Email Address: Fax Number:	
Additional User Information – Responsible for pa	erforming eBooking operation.
Login ID: Full Name: Phone Number: Email Address: Fax Number:	
Additional User Information – Responsible for pe	erforming eBooking operation.
Login ID: Full Name: Phone Number: Email Address: Fax Number:	

Private and Confidential – For Internal Use Only



GLOBAL SOURCING DEPARTMENT

300 Phillipi Road P.O. Box 28512 Columbus, Ohio 43228-0512 U.S.A. Phone (614) 278-6800

Fax (614) 278-6475

IMPORTER'S BLANKET STATEMENT of NON-REIMBURSEMENT ANTI- DUMPING DUTIES

Effective Date:	Immediate
Importer of Record:	Big Lots Stores, Inc. 300 Phillipi Road Columbus, OH 43228
I.R.S:	#31-1186811-00
Manufacturer:	Various
Antidumping Case:	
manufacturer, producer, seller or expo from China, which have been and/or w	
· · · · · · · · · · · · · · · · · · ·	Service will be notified if there is any reimbursement of dumping duties by the rter to the importing company at any time in the future.
Failure to file this certificate prior to li anti-dumping duties.	quidation will result in the presumption of reimbursement and assessment of double

Big Lots Stores, Inc

VP Global Sourcing

STATEMENT REGARDING THE IMPORTATION OF RADIO FREQUENCY DEVICES CAPABLE OF CAUSING HARMFUL INTERFERENCE

(Read instructions before completing form. Please type or print clearly in ink.)

Part I - All Blocks	MUST Be Co	mpleted	77		3/2			
Date of Entry	Entry Numb	er Po	rt of Entry ¹	Harm	onized Tariff Num	ber ²	Quantity of Item (not	number of containers)3
Device Model/Type	e Name or #	Trade	Name	F	CC ID		Description of E	Equipment
Manufacturer's	Name and Ado	dress	Cor	isignee's N	lame and Address		Importer's Name and	Address
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Printed or Typed N	ame of Imports	er or Cons	ignee		Signature of Imp	orter o	or Consignee	Date (Month/Day/Year)
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Warning: Any person who knowingly makes a false declaration may be fined not more than \$250,000 or imprisoned not more than 5 years, or both, pursuant to 18 U.S.C. \$1001.

Part II - With Regard to the Importation of the Described Radio Frequency Device(s), I DECLARE THAT: (Place an "X" in only one box)
1. The FCC has issued a grant of equipment authorization for the FCC ID listed above.
2. An FCC grant of equipment authorization and an FCC ID are not required, but the equipment complies with FCC technical requirements.
3. The described equipment is being imported in limited quantities for testing and evaluation for compliance with technical requirements or marketing suitability. The equipment will not be offered for sale or otherwise marketed. (See Instructions)
4. The described equipment is being imported in limited quantities for demonstration at industry trade shows and will not be offered for sale or otherwise marketed. (See Instructions)
5. The described equipment is being imported solely for export. It will not be offered for sale or otherwise marketed in the U.S.
5(a). The described equipment is a non-U.S. standard cellular phone that can only function outside of the U.S. (See Instructions)
6. The described equipment is being imported for use exclusively by the U.S. Government.
7. Three or fewer radio receivers, computers, or other unintentional radiators as defined in Part 15 of the FCC Rules, are being imported for an individual's personal use and are not intended for sale.
8. The described equipment is being imported for repair and will not be offered for sale or otherwise marketed.

- Port of Entry Use Schedule D Classification of U.S. Customs Districts and Ports for U.S. Foreign Trade Statistics a four digit code i.e., New York City, NY 1001.

 Harmonized Tariff Number Harmonized Tariff Schedule of the United States.

 This quantity must be total number of items, not number of containers.

FCC Form 740 March 2004

INSTRUCTIONS FOR COMPLETION OF FCC FORM 740

This form must be completed for each radio frequency device, as defined in 47 U.S.C. 302 and 47 C.F.R. 2.801, which is imported into the Customs territory of the United States. The original shall be filed with the U.S Customs Service on or before the date the shipment is delivered to a U.S. port of entry.

The completed form must accompany each such entry.

The following are typical examples of devices that require the use of FCC Form 740: radio and TV receivers, converters, transmitters, transmitting devices, radio frequency amplifiers, microwave ovens, industrial heaters, ultrasonic equipment, transceivers, and computers.

Marketing, as used in this form (and 47 C.F.R. 2.1201 et seq.), means sale or lease (including advertising for sale or lease, or display at a trade show) or import, ship or distribute for the purpose of selling or leasing or offering for sale or lease.

Limited quantities, as used in this form, are the number specified in 47 C.F.R. 2.1204(a)(3). Waivers of this limit are infrequently granted but may be requested from the FCC office listed in 47 C.F.R. 2.1204(a)(3)(iii). Written waiver requests must contain specific information required by that office.

Equipment imported for test, evaluation or display (see import conditions 3 or 4 of Part II of this form) may not be marketed (sold or leased, offered for sale or lease, advertised, etc.). Display of this equipment must include markings clearly indicating that the device(s) are not eligible for sale. See 47 C.F.R. 2.803 for details regarding this labeling.

Wireless telephony devices that do not have a FCC grant of equipment authorization must either comply with 47 C.F.R. 2.1204(a)(5) or 47 C.F.R. 2.803(a)(2) (e.g., Verification or Declaration of Conformity is required).

The identification (company name and model number/FCC ID) of the radio frequency device specified on the front of this form must be identical to the company name and model number/FCC ID inscribed on the device. If the device being imported requires an equipment authorization to be issued by the FCC (e.g., Certification), it is important that the name of the company, description of the device and FCC ID specified on the grant of equipment authorization agree exactly with the same information shown on the front of this form. Any discrepancy between the information on this form and the FCC grant of equipment authorization may result in unnecessary delays, additional expense, or enforcement action.

FCC Form 740 may be reproduced provided the following conditions are met (see 47 C.F.R. 0.409, Commission Policy on Private Printing of FCC Forms.) Some of the conditions are listed below:

- That private companies reproducing the form use a printing process resulting in a product that is comparable to the original document;
- That private companies reproducing the form refrain from including therein or attaching thereto any advertising matter or deleting any material from the form;
- 3. That private companies reproducing the form exercise care that the form being reproduced or distributed is the current edition presently used by the FCC for the type of application involved: such private company to be advised that, though the Commission will endeavor to keep the public advised of revisions of the form, it cannot assume responsibility to the extent of eliminating any element or risk against overstocking, etc.

PAPERWORK REDUCTION ACT STATEMENT AND PRIVACY ACT STATEMENT

The solicitation of information requested on this form is authorized by the Communications Act of 1934, as amended. The information collected will be used to ascertain whether equipment authorization is required, and if so, whether or not it has been granted. If all the information is not provided the importation of this or other shipments may be delayed or prevented. Accordingly, every effort should be made to provide all necessary information. Your response is required to obtain a benefit.

Public reporting for this collection of information is estimated to average .04 seconds per response, including the time for reviewing instructions searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing the burden, should be sent to the Federal Communications Commission, Performance and Evaluations and Records Management, Washington, DC 20554, Paperwork Reduction Project (3060-0059) DO NOT SEND COMPLETED FORMS TO THIS ADDRESS. Individuals are not required to respond to a collection of information unless it displays a currently valid OMB control number.

THE FOREGOING NOTICE IS REQUIRED BY THE PRIVACY ACT OF 1974, P.L. 93-579, DECEMBER 31, 1974, 5 U.S.C. 552A(E)(3), AND THE PAPERWORK REDUCTION ACT OF 1995, P.L. 104-13, OCTOBER 1, 1995, 44 U.S.C. 3507. FCC Form 740 instructions – Page 2

DEPARTMENT OF HEALTH AND H	UMAN SERVICES	Form Approved OMB No. 0910-0025 Expiration Date: 11/30/2003	WWW of the second
PUBLIC HEALTH SER FOOD AND DRUG ADMINIS DECLARATION FOR I ELECTRONIC PRODUCTS RADIATION CONTROL S	MPORTED SUBJECT TO		less requested. is, submit the following to FDA: n (e.g. CF 3461, CF 3461 Alt,
U.S. CUSTOMS PORT OF ENTRY		ENTRY NUMBER	DATE OF ENTRY
NAME & ADDRESS OF MANUFACTURING SITE;	COUNTRY OF ORIGIN	NAME & ADDRESS OF IMPORTER &	ULTIMATE CONSIGNEE (if not importer)
PRODUCT DESCRIPTION	QUANTITY (Items/Containers		
DECLARATION: 1 / WE DECLARE THAT TH			atements, fill in blanks, & sign)
☐ A. ARE NOT SUBJECT TO RADIATION			
1. Were manufactured prior to the effecti 2. Are excluded by the applicability claus Specify reason for exclusion	ve date of any applicable star e or definition in the standard	or by FDA written guidance.	
 3. Are personal household goods of an in 4. Are property of a party residing outsidents. 5. Are components or subassemblies to 6. Are prototypes intended for on going present of the prototype. 7. Are being reprocessed in accordance or transferred without FDA approval. 	e the U.S. and will be returned be used in manufacturing or a product development by the in e., not distributed). (Quantitie	I to the owner after repair or servicing. Is replacement parts (NOT APPLICABLE to porting firm, are labeled "FOR TEST/EVAL Is Limited - see reverse.)	diagnostic x-ray parts). .UATION ONLY," and will be exported,
1. Last annual report or Product/Initial re	TO THIS EFFECT IS AF	FIXED TO EACH PRODUCT. COM	PLIANCE DOCUMENTED IN:
ACCESSION NUMBER o		MANUFACTURER OF RECORD (Filed rep	ort with FDA/CDRH)
C. DO NOT COMPLY WITH PERFOR	RMANCE STANDARDS; RCE; WILL BE USED UN TOMS SUPERVISION V Training (attach Form FDA 760	DER A RADIATION PROTECTION F WHEN THE FOLLOWING MISSION I	PLAN; AND WILL BE DESTROYED
D. DO NOT COMPLY WITH PERFORMINTRODUCED INTO COMMERCE INTO COMPLIANCE IN ACCORDATION 1. Approved Petition is attached.	UNTIL NOTIFICATION	S RECEIVED FROM FDA THAT PR PROVED PETITION. (See Form FD.	ODUCTS HAVE BEEN BROUGHT
WARNING: Any person who knowingly declaration may be fined not more the imprisoned not more than 5 years or both, 18 U.S.C. 1991.	an \$10,000 or pursuant to Title	RE OF IMPORTER OF RECORD	5
18 U.S.C. 1001. Any person importing a electronic product may also be subject to a \$1000 per violation, up to a maximum \$300 violations pursuant to Title 21 U.S.C. 360pp.	civil penalties of	D TITLE OF RESPONSIBLE PERSON	
Public reporting burden for this collection of searching existing data sources, gathering and me this burden estimate or any other aspect of this compared from the food and Drug Administ CDRH (HFZ-342) 2094 Gaither Road Rockville, MD 20850 An agency may not conduct or sponsor, and a performance of the food from the food	naintaining the data needed, a ollection of information, includ tration	nd completing reviewing the collection of info ng suggestions for reducing this burden to:	rmation. Send comments regarding
FORM FDA 2877 (12/00)	PREVIOUS ED	TION IS OBSOLETE CONTRACTOR	PAGE 1 OF 2 PAGES ER

INSTRUCTIONS TO IMPORTERS/BROKERS OF ELECTRONIC PRODUCTS

PURPOSE: The Form FDA 2877 must be completed for electronic products subject to Radiation Control Standards (21 CFR 1010 and 1020-1050) prior to entry into the United States. The local Food and Drug Administration (FDA) district office will review the declaration and notify the importer/agent if the products may be released into U.S. commerce or if they must be held under bond until exported, destroyed, or reconditioned. Until the shipment is released, it may be subject to redelivery for FDA examination.

PAPER OR ELECTRONIC SUBMISSION: Paper entries may be made by submitting the signed original FDA 2877 along with U.S. Customs forms to the local FDA district office; if electronic products are given a MAY PROCEED, a signed copy of CF 3461 will be returned, or if not given a MAY PROCEED, a FDA Notice of Action will be issued. For electronic entries, follow U.S. Customs Service ACS/ABI format and procedures, supported by a signed copy of this form or similar letter. Multiple entries of the same product and model families that are filed electronically may be supported by one form dated not more than 12 months previously.

DECLARATION: Select A, B, C, or D and then select the appropriate number; fill in requested information and sign. For electronic entries, AofC (affirmation of compliance) = RA#, RB#, RC#, or RD# (e.g., Radiation Declaration A5 = RA5). Transmit model number using AofC code MDL and transmit brand name using FDA line level brand name field. If RA3 or RA6 is selected, you must transmit quantity (number of units) using the Quantity and Unit of Measure Pairs at the FDA line level.

DECLARATION A: Importers should be prepared to demonstrate compliance to or non-applicability of FDA standards, regulations, or guidance. Components or sub-assemblies must be non-functioning. Products being reprocessed must be exported by the importer, without intermediate transfer of ownership. For RA3 the quantity limit is 3 and for RA6 the limit = 50 units TV products, microwave ovens, and Class 1 laser products limit = 200 units CD-ROM and DVD (digital versatile disc) laser products; see May 14, 1997, notice to industry issued by the Center for Devices and Radiological Health (CDRH).

DECLARATION B: If declaration RB1 is selected, provide the FDA Establishment Identifier (FEI) of the manufacturer who filed the radiation product/abbreviated report to FDA, CDRH, Rockville, Maryland. To transmit the accession number of that report use AofC code ACC. If the manufacturer cannot be determined or located, the importer must be able to provide evidence showing a certification (certifi.) label on each product and state reason: returned to orig exporter or certifi. label evidence. The new AofC codes (RB1, RB2) for this declaration will not be activated until a process is made available to determine the FEI of the responsible firm. Continue to use RAB in electronic transmission until the FEI query is available and industry is notified of its availability.

DECLARATION C: Noncompliant products may be imported only for research, investigations/studies, demonstration or training. They should be used only by trained personnel and under controlled conditions to avoid unnecessary radiation exposure. Product(s) will be detailed by the local FDA district office. Since product(s) for which "C" Declarations are made will be under Temporary Import Bond (TIB) or equivalent, ultimate disposition is limited to export or destruction under U.S. Customs supervision when the purpose has been achieved or the length of time stated has expired. For purposes other than demonstration, the Form FDA 766, outlining protections, must be approved by FDA prior to use. The importer/broker must include with the FDA 766:

- A full description of the subject electronic product(s). The purpose for which the product(s) is being imported.
- How the product(s) will be used.
 Where the product(s) will be located.
- The approximate length of time and dates the product(s) will be in this country.

For product(s) being used for trade shows/demonstrations, list the dates and use restrictions (Form FDA 766 is not required). A sign stating that the product does not comply with FDA performance standards must be displayed and viewable at all times during the use of product(s). All medical products, cabinet x-ray, or Class IIIb and IV lasers may NOT operate (turn on product(s)) at trade shows.

DECLARATION D: Noncompliant products must be brought into compliance with standards under FDA supervision and following a plan approved by FDA. The plan, documented on the Form FDA 766, must address technical requirements, labeling, and reporting. Some plans may need approval by both the CDRH and the local FDA district office. Use of this declaration is limited to occasional shipments; ongoing reconditioning is considered manufacturing that is handled through other means. Product(s) will be detained by the local FDA district office. An FDA 766 must be filed indicating the procedure intended to bring the product into compliance. This procedure will include a satisfactory corrective action plan and/or a product report. The FDA 766 must include all of the information requested under Declaration C. The approximate length of time will be for the amount of time needed to bring product(s) into compliance. Declaration D is also made for failure to provide reports, failure to certify, etc.

If an importer/broker intends to import equipment into the United States for purposes of research, investigation, studies, demonstrations, or training but also wishes to retain the option of bringing the product into compliance with the performance standard, check Declarations C and D on the FDA 2877 and insert the word "or " between the Affirmations. Note: The U.S. Customs Service will treat this entry as a "D" Declaration for purposes of duty. Such requests must be made on the FDA 766; include Items 1, 2, and 3 under Declaration C, a statement of the need to use the option "C" or "D" Declaration, a statement of how the product(s) will be brought into compliance and the approximate length of time necessary to evaluate or demonstrate the product(s) and the time necessary to bring the product(s) into compliance (both actions must be accomplished within the period of time granted by FDA). For electronic entries select Declaration RD3.

Ultimately, product(s) must be brought into compliance with the applicable standard in accordance with a corrective action plan which has been approved by the FDA. If the product(s) are not brought into compliance within the allotted time frame of the approved application and an extension is not requested of, or granted by, the FDA, the local FDA district office shall refuse entry on the shipment and require the product(s) to be either exported or destroyed under U.S. Customs supervision.

If additional guidance is needed, please contact your local FDA district office or consult the following FDA web pages: www.fda.gov/cdrh, www.fda.gov/ora/hier/ora_field_names.txt, and www.fda.gov/ora/compliance_ref/rpm_new2/contens.html.

[Ref: 21 U.S.C. 360mm, 21 CFR 1005, 19 CFR 12.90-12.91.]

FDA: CP 7382.007/.007A

FORM FDA 2877 (12/00)

PREVIOUS EDITION IS OBSOLETE.

PAGE 2 OF 2 PAGES

	SERVICES	EXPIRATION DATE: April 30, 2008
FOOD AND DRUG ADMINISTRATION REGISTRATION OF DEVICE ESTABLISHMENT	ABLISHMENT	R VALIDATION (FDA USE ONLY)
Submit an original copy, Please do not mail the instruction pages with your form (this page only). Return form to: Prood and Dury Administration Center for Devices & Radiological Health, HFZ- 308 9200 Corporate BIVd. Rock	6. REASON FOR UPDATE (check all that apply) 6. Establishment Name Change 6.2 Establishment Type Change (deletion or addition) 6.2 Establishment Address Change - Merged with Other 6.3 Establishment Address Change - Merged with Other	(0)
3. TODAY'S DATE (mm/dd/yyy) 07/03/06	6.4	T.3 Foreign Exporter T.4 Initial Distributor/Importer
4. TYPE OF REGISTRATION 4.1 Initial 4.2 Update 4.3 Preproduction	Conficial Correspondent Name/Address Change	
5. REQUIRED TO SUBMIT DEVICE LISTING (Form FDA 2892)? [] Yes [] No; If No, Explain:		7.7 Repackager/Relabeler 7.8 Reprocessor of Single Use Devices 7.9 Specification Developer 7.10 U.S. Manufacturer of Export Only Devices
8. ESTABLISHMENT (No P.O. Boxes) Business Name	6.10 No Longer a Device Establishment 6.11 In Production 6.12 Trade Name or Establishment URL Change	10. U.S. AGENT NAME AND ADDRESS (Foreign Establishments Only) Same as Official Correspondent? [IN No. List Individual's Name, Title, Business Name, Number & Street, City, State, 219 Code, Nov D. Boxes, The U.S. Agent must either reside in the IS or maintain a place of business there.
Number & Street	9. OFFICIAL CORNES TOWNERS I mannages sequined; Reason for OC Name Change (see instructions):	Name
City State ZIP Code	Name	Title
Foreign State Postal Code Country	Business Name	Business Name
11. OWNEP/OPERATOR (No P.O. Boxes) Rusineses Name	Number & Street	Number & Street
O I VANIE OF THE PROPERTY OF T	City State ZIP Code	City
Number & Street	Foreign State Postal Code Country	State ZIP Code
City State ZIP Code	9.1 EMAIL	10.1 EMAIL
Foreign State Postal Code Country	9.2 PHONE NO. (Phone no. should include area code or country/city codes)	10.2 PHONE NO. IN U.S. (Phone no. should include area code)
11.1 PHONE NO. (Phone no. should include area code or country/city codes)	es) 9.3 FAX NO. (FAX no. should include area code or country/city codes)	10.3 FAX NO. IN U.S. (FAX no. should include area code)
12. OTHER BUSINESS TRADING NAMES		
13. SIGNATURE OF OFFICIAL CORRESPONDENT	13.1 PRINTED NAME (Mr., Miss., Ms., Ms., Dr.)	13.2 ПП.Е
14. ESTABLISHMENT'S URL (Optional):		
NOTE: This form is authorized by Section 510 of the Federal Food, Drug, and U.S.C.331(p). Persons who violate this provision may, if convicted, be subject respect is a violation of Section 301(p)(2), (21, S.C. 331(p)), and may be a virtually unlied to a virtually unlied of the section of th	NOTE: This form is authorized by Section 510 of the Federal Food, Drug, and Ossmetic Act (21 U.S.C. 360). Failure to report this information is a violation of Section 301(p) of the Act (21 U.S.C. 361(p)). Persons who violate this provision may, if convicted, be subject to a fine or imprisonment or both. The submission of any report that is false or misleading in any material respect is a violation of Section 301(q)(s) and many be a vial many be a vial and any and a person is not required to respond to a necessary as a currently valid. Other contribution in the violation of section 301(q)(s) and Act of the provision of the Act (21 U.S.C. 1001).	n 301(p) of the Act (21 acading in any material aquired to respond to a O / W / P / Y

Instructions for Completing FDA Form 2891: Registration of Device Establishment - All information submitted must be in English. Submit a signed original copy.

Registration Number - Fill in if a registration number has been previously assigned by the Food a Administration (FDA). Leave this space blank if no registration number has been issued. FDA assign a registration number after processing and provide this to the official correspondent. s been assign Item 2. Owner/Operator Number - Fill in if an owner/operator Identification (I.D.) number has previously state by the Foresting this space ablank fin oil. number has been issued. FDA will as an I.D. number has been issued. FDA will as an I.D. number processing and provide this to the official correspondent.

Item 3. Today's date - Enter the month, day, and year the form is completed using a MM/DD/YYYY date format.

Item 4. Type of registration:

4.1 Initial - Check this if it is the first time you are registering this establishment. If you are required to establishment, the United registration form must be submitted with the initial isting form, and in the case of a foreign establishment, the United States agent information (flem #10). If the information is not submitted together, the initial registration form will not be accepted.

4.2 Update - Check this box if the form is being submitted to indicate changes in registration information that you have already submitted to the FDA. Only fill in the section/items that have changed and items 1 and 2.

4.3 Preproduction - Check this if you will not start producing medical devices for at least 3 months. You must notify DA white you start producting medical devices. If the establishment does not notify the Ventre fro Devices and Radiological Health (CDRH) that it has begun an activity that requires registration within 6 months, the preproduction registration form will then be archived without further processing. You are not officially registered until you have notified FDA of your active status by submitting an update FDA 2891, and checking box 6.11, In Production.

Item 5. Required to Submit Listing - Check Yes or No. If No, explain why you are not required to list.

Who Must List - An owner/operator of an establishment not exempt under 21 CFR 807.65 who is engaged in the manufacture, preparation, propagation, compounding, assembly or processing of a medical device intended for commercial distribution (marketing) is required to list its device on form FDA 2892 within 30 days of entering the device into commercial distribution in the U.S. This includes manufacturers, repeakagers and featbelies; specification developers, reprocessors of single-use devices, remanufacturers, foreign exporters and U.S. manufacturers of export only devices.

Domestic contract manufacturers and sterifizers that commercially distribute their devices must register and start their devices. Except in unique circumstances, initial distributors are the only establishment type not required to list.

Item 6. Reason for Update - Only fill this item out if type of registration is Update. Check all that apply

6.1 Establishment Name Change - A change in the establishment name

6.2 Establishment Type Change - A change in the types of activities which require registration as a medical device establishment (see Item 7 - Establishment Types).

6.3 Establishment Address Change - Werged with Other Establishment - Indicates establishment has merged with another establishment, and the new company is located at a different address from this registration.

6.4 Establishment Address Change - Moved to New Location - Change in the establishment address due only to a physical relocation

6.5 Official Correspondent Name/Address Change - Any change in the Official Correspondent

nformation, including name and address.

6.6 U.S. Agent Change - Any change in the US Agent information, including name and address.

6.7 Owner/Operator Name/Address Change - Same Company New Name or Address - Indicates only a new name and address for Owner/Operator, but establishment remains under same ownership.

6.8 Owner/Operator Change - Sold Establishment - Indicates Owner/Operator has changed because establishment sold to another firm.

6.9 Out of Business - The establishment has ceased to exist as an identifiable organization.

6.10 No Longer a Device Establishment - The establishment is no longer engaged in activities which require it to be registered as a medical device establishment, but the establishment is still in existence for other activities or purposes.

FORM FDA 2891 (7/05)

6.11 In Production - Notification that an establishment has gone from a "pre-production" status to an "in production" status.

6.12 Trade Name or Establishment URL Change - Any change in the Other Business Trading Names or the Establishment's URL. Item 7. Establishment Types - You should only select the establishment types that apply to to operations performed at the establishment you are registering. Check all that apply to the establishment.

7.1 Contract Manufacturer - Manufactures a finished device to another establishment's specifications and puts device in commercial distribution.

7.2 Contract Sterilizer - Provides a sterilization service for another establishment's devices and puts the devices in commercial distribution.

7.3 Foreign Exporter - Exports or offers for export to the United States (U.S.), a device manufactured or processed by another individual, partnership, corporation or association in a foreign country, including devices originally manufactured in the United States. A foreign exporter must have an establishment address outside the U.S.

7.4 Initial Distributor/Importer - Takes first title to devices imported into the U.S. An Initial Distributor must have a U.S. address.

7.5 Manufacturer - Makes by chemical, physical, biological or other procedures, any article that meets the definition of "device" in Section 201(h) of the Federal Food, Drug, and Cosmetic (FD&C) Act.

7.6 Remanufacturer - Processes, conditions, renovates, restores, or does any other act to a finished device that significantly changes the finished device's performance or safety specifications, or in any way obtanges the literated use.

þ Relabeler - Changes the content of the labeling from that supplied from the original manufacturer is distribution under the establishments own name. A relabeler does not include establishments that do rehange the original labeling but merely add their own name. 7.7 Repackager - Packages finished devices from bulk or repackages devices made manufacturer into different containers (excluding shipping containers).

7.8 Reprocessor of Single Use Devices - Performs remanufacturing operations on a single use device.

7.9 Specification Developer - Develops specifications for a device that is distributed under the establishment's own name but performs no manufacturing operations on the device. This includes establishments that in addition to developing specifications also arrange for the manufacturing of devices labeled with another establishment's name by a contract manufacturer.

7.10 U.S. Manufacturer of Export Only Devices - Manufactures medical devices that are not sold in the U.S. and are manufactured solely for export to foreign countries.

Item 8. Establishment Name and Address:

Name - Enter the legal name of the establishment conducting the regulated activity

Address - Number and Street - Enter the number and street at which the registering establishment is located. Do NOT use postal box or rural route numbers.

Domestic Establishments:

City - Enter the city in which the establishment is located.

State - Enter the two-character state code of the U.S. Postal Service for the state, territory, or pos-

ZIP Code +4 - Enter the U.S. postal ZIP code +4 (if known)

Foreign Establishments:

Foreign State - Enter the foreign state (i.e., province, prefecture, region, territory) names in which the sstablishment is located.

Postal Code - Enter the foreign country postal code

Foreign Country Name - Enter the full foreign country name.

(Continued on next page)

Instructions for Completing FDA Form 2891: Registration of Device Establishment - All information submitted must be in English. Submit a signed original copy.

(Continued from previous page)

Item 9. Official Correspondent Name and Address:

Reason for OC Name Change - Briefly state why the OC Name is changing, e.g., OC left company, OC deceased, new owner had changed OC, OC moved onto new position in company.

Name - Enter the name of the individual designated as the official correspondent for registration and **Business Name -** Enter the name of the establishment, owner or operator, or other place of business, with which the official correspondent is associated. isting purposes

Address - Number and Street - Enter the number and street or post office box of the official correspondent's place of business. A post office box number is acceptable.

Domestic Correspondents:

City - Enter the city in which the official correspondent is located

9 State - Enter the two-character state code of the U.S. Postal Service for the state, territory,

ZIP Code +4 - Enter the U.S. postal ZIP Code +4 (if known).

Foreign Correspondents

Foreign State - Enter the foreign state (i.e., province, prefecture, region, territory) names in which the establishment is located.

Postal Code - Enter the foreign country postal code.

Foreign Country - Enter the full foreign country name

9.1 Email - Enter the email address of the official correspondent (NO generic accounts, must be OC's

9.2 Phone Number - Include country, city, area code, number and extension

9.3 Fax Number - Include country, city, area code, and number.

Item 10. U.S. Agent Name and Address - The United States agent must either reside in the U.S. or maintains a place of business in the U.S. If the U.S. agent is also the Official Correspondent, check yes and you do not need to provide the address information but make sure that there is a U.S. phone and fax number. If the U.S. agent is a different person then check no and fill in the following information:

Individual's Name - Enter the name of the individual designated as the U.S. agent

Agent's Title - Print/type the title of the U.S. agent

Business Name - Enter the name of the place of business with which the U.S. agent is associated

Number and Street - Enter the number and street of the U.S. agents's place of business. A post office box number is NOT acceptable.

City - Enter the city in which the U.S. agent is located.

State - Enter the two-character state code of the U.S. Postal Service for the state, territory,

ZIP Code +4 - Enter the U.S. postal ZIP Code +4 (if known)

Public reporting burden for this collection of information is estimated to average .25 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to the address to the right.

3ockville, MD 20850-4015 9200 Corporate Blvd

An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number.

FORM FDA 2891 (7/05)

10.1 Email - Enter the email address of the U.S. agent

10.2 Phone Number In U.S. - Include area code, number and extension.

10.3 Fax Number in U.S. - Include area code, number and extension.

Item 11. Owner/Operator Name and Address:

is the owner or operator of the registering establishment. Only enter the proprietor's name or an individual's name if no other business name exists (abbreviate only if necessary). Name - Enter the business name of the corporation, subsidiary, affiliated company, or partnership that

Address - Number and Street - Enter the number and street at which the Owner/Operator is located. Do NOT use postal box or rural route numbers.

Domestic Owner/Operators:

City - Enter the city in which the owner/operator is located.

State - Enter the two-character state code of the U.S. Postal Service for the state, territory, or pos-

ZIP Code +4 - Enter the U.S. postal ZIP Code +4 (if known)

Foreign Owner/Operators:

Foreign State - Enter the foreign state (i.e., province, prefecture, region, territory) names in which the owner/operator is located.

Postal Code - Enter the foreign country postal code.

Foreign Country - Enter the full foreign country name.

11.1 Phone Number - Include country, city, area code, number and extension

Item 12. Other Business Trading Names - Enter any other establishment names used. Please put a semi-colon between the names. Do not include the names of distributors for whom this establishment makes devices. Do not list registered trademarks in use by the firm.

Item 13. Signature of Official Correspondent - The signature of the designated official correspondent.

13.1 Name - Print/type the name of the official correspondent.

13.2 Title - Print/type the title of the official correspondent.

Item 14. Establishment's URL (Optional) - List the main web address associated with this establishment.

Center for Devices and Radiological Health (HFZ-308)

Food and Drug Administration

NOTE: This form is authorized by Section 510 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360), Failure to report this information is a violation of Section 301(p) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301(p)). Persons who violate this provision may, if concluded, be subject to a fine or imprisonment or both. The submission of any place or infederal respect is a violation of Section 301(p)(2), (21 U.S.C. 331(q))(2)) and may be provision may, if a violation of Section 301(p)(2), (21 U.S.C. 331(q))(2)) and may be a violation of 18 U.S.C. 1001. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information mines it displays a currently valid OMB control number. 2. OWNNER/OPERATOR NAME (Business name) 3. OWNNER/OPERATOR NAME (Business name) 3. OWNNER/OPERATOR NAME (Business name) 6. ESTABLISHMENT NAME (Business name) 7. ESTABLISHMENT NAME (Business name) 8. CITY 8. CODE 8. CITY 8. CODE 8. CITY 9. CODE 9. COUNTRY FOREIGN STATE 10. C. C331(q)(2), Persons who violate this provision may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)) and may be avoidable to solicit of the Act (31 U.S.C. 301(q)(2)) and may be avoidable to solicit of the Act (21 U.S.C. 331(q)(2)	ug, and Cosmetic Act (21 U.S.C. 360), Failure to rep	in a all mailtonness and will all the second of an	* * * * * * * * * * * * * * * * * * * *	
2. OWNER/OPERATOR NUMBER 3. OWNER/OPERATOR NAME (Business name) NUMBER AND STREET CITY FOREIGN STATE FOREIGN STATE	both. The submission of any report that it, and a person is not required to respon	is false or misleading in any mater to a collection of information unles	olation of Section 301(p) of ial respect is a violation of se it displays a currently va	f the Act (21 U.S.C.331(p)). Persons who Section 301(q)(2), (21 U.S.C. 331(q)(2)) iid OMB control number.
WINER/OPERATOR NAME (Business name) BER AND STREET FIGN STATE		4. REGISTRATION NUMBER		
BER AND STREET POSTAL CODE		5. ESTABLISHMENTNAME (Business name)	siness name)	
EIGN STATE POSTAL CODE		NUMBER AND STREET		
	ZIP CODE	СІТУ		STATE ZIP CODE
	COUNTRY	FOREIGN STATE	POSTAL CODE	COUNTRY
6. LISTING INFORMATION: Number of product codes you are going to list for this establishment:	for this establishment:			
REASON FOR LISTING: New Listing Update to Device Already Listed	ready Listed Delete Listing	REASON FOR LISTING:	New Listing Upo	Update to Device Already Listed Delete Listing
PRODUCT CODE PMA NUMBER	510(k) NUMBER	PRODUCT CODE	PMA NUMBER	510(k) NUMBER
CLASSIFICATION NAME		CLASSIFICATION NAME		
PROPRIETARY NAME		PROPRIETARY NAME		
COMMON OR USUAL NAME		COMMON OR USUAL NAME		
PREVIOUS LISTING NUMBER	B E442476	PREVIOUS LISTING NUMBER		LISTING NUMBER E442477
Contract Manufacturer Remanufacturer Specific Contract Sterilizer Remanufacturer Specific Poreign Exporter Bepackager/Relabeler U.S. Ma	Reprocessor of Single Use Devices Specification Developer U.S. Manufacturer of Export Only Devices	Contract Manufacturer Contract Sterilizer Foreign Exporter	Manufacturer Remanufacturer Repackager/Relabeler	Reprocessor of Single-use device Specification Developer U.S. Manufacturer of Export Only Devices
REASON FOR LISTING: New Listing Update to Device Already Listed	Iready Listed Delete Listing	REASON FOR LISTING:	New Listing Upo	Update to Device Already Listed
PRODUCT CODE PMA NUMBER	510(k) NUMBER	PRODUCT CODE	PMA NUMBER	510(k) NUMBER
CLASSIFICATION NAME		CLASSIFICATION NAME		
PROPRIETARY NAME		PROPRIETARY NAME		
COMMON OR USUAL NAME		COMMON OR USUAL NAME		
PREVIOUS LISTING NUMBER	R E442478	PREVIOUS LISTING NUMBER		LISTING NUMBER E442479
Contract Manufacturer Remanufacturer Specific Contract Sterilizer Remanufacturer Specific Foreign Exporter Repackager/Relabeler U.S. Ma	Reprocessor of Single Use Devices Specification Developer U.S. Manufacturer of Export Only Devices	Contract Manufacturer Contract Sterilizer Foreign Exporter	Manufacturer Remanufacturer Repackager/Relabeler	Reprocessor of Single Use Devices Specification Developer U.S. Manufacturer of Export Only Devices
7. SIGNATURE OF OFFICIAL CORRESPONDENT	8. TYPED OR PRINTED NAME		ПТСЕ	

Instructions for Completing FDA Form 2892: Device Listing All information must be in English. Submit a signed original copy.

Item 1, Today's date - Enter the month, day, and year the form is completed using a MM/DD/YYYY date format.

Item 2. Owner/Operator Number - Fill in if an owner/operator Identification (I.D.) number has been previously issued by the FDA. Leave this space blank if no I.D. number has been issued. FDA will assign an I.D. number after processing and provide this to the official correspondent.

Item 3. Owner/Operator Name & Address - Name - Enter the business name of the corporation, subsidiary, affiliated company, or partnership that is the owner or operator of the registering establishment. Only enter the proprietor's name or an individual's name if no other business name exists. Address - See General Address instructions

Item 4. Registration Number - Fill in if a registration number has been previously assigned by the Food and Drug Administration (FDA). Leave this space blank in registration mumber has been issued. FDA will assign a registration number after processing and provide this to the official correspondent.

Item 5. Establishment Name and Address - Name - Enter the legal name of the establishment conducting the regulated activity. Address - See General Address instructions.

tem 6. Number of product(s) that you are going to list for this establishment: Enler the number of product codes that you are now listing on this form. For each of the product codes fill in the following information:

Reason for the Listing: Check the appropriate box. Select only one.

Product Code: Enter the three letter product code that corresponds to the device name assigned to your delvero, or enter the three letter product code that appears on the STO(k) clearance or PMA approval eleter. The device names and product codes appear in an on-line database, www.rda.gov/cdr/prodcode-hml. If you cannot determine the product code, provide the STO(k), PMA or regulation number of the device. Do not confuse the product code with the seven digit regulation number assigned to the type of device classified in the Code of Federal Regulation. Fars 862-892.

All device types classified as exempt from the 510(k) requirements are subject to the limitations of exemptions of device examplions are found in the device classification chapters in 21 CFB xxxx, y, where xxx is replaced with Parts 862-892 (e.g., 862-9, 864-9, etc.). It is your responsibility to ensure that you may be an exceeded the limitations of exemption. If your device does not exceed the limitations of exemption, your device acceeds the limitations of exemption, you must submit a 510(k) and receive a letter from FDA staining that your device may be commercially distributed in the U.S. prior to marketing your device.

PMA Number: Please enter the PMA number found on your Premarket Approval (PMA) letter.

510(k) Number: Please enter the 510(k) number found on your Substantial Equivalency (SE) letter.

Classification Name: Enter the classification name or device name for the generic category of the device. The name can be found in the on-line product classification database, www.tda.gov/prodoce/html. DO NOT ENTER "NONE" OR "WILLITPLE". IN THIS BLOCK. If you are unable to determine your classification name, please leave this item blank and submit a copy of the device labeling and an explanation of the intended use of the product. If you know the name of a similar product, please also provide this information. FDA staff will review the labeling and an intended use information and either determine the correct product code, or advise you of any further actions you may need to take.

Proprietary Name: Multiple brand names should be entered. FDA does not issue a separate medical device listing number for each brand name of your device.

Common or Usual Name: If more than one device is being listed under one classification name, enter a descriptive phrase which represents the group of devices, i.e., "Types of Broguers," or "X-ray Systems." If one or more devices represented by a classification name is labeled and marketed as "sterile", include the word sterile as part of the common or usual mame, i.e., 'Sterile and Non-Sterile Syringes."

Pervious Listing Number: If this is an update to a previously listed device, please enter the Document Number that appears on the last Device Listing form you submitted for the same product code and establishment registration number.

Establishment Types: Check all that apply. You should only select the establishment type(s) that apply to the operations performed at the establishment you are registering. For example, if the establishment is a manufacture and specifications developer, but only manufactures the listing device then only check. nanufacturer

Please make sure that the codes you select match the establishment type(s) that are on file with FDA in the establishment's registration data. If FDA finds a mismatch, your form will not be processed until you update the establishment's registration data. The listing establishment types are defined as follows.

Contract Manufacturer - Manufactures a finished device to another establishment's specifications and puts device in commercial distribution. Contract Sterilizer - Provides a sterilization service for another establishment's devices and puts the devices in commercial distribution.

processed by another individual, partnership, corporation or association in a foreign country, including devices originally manufactured in the United States. A foreign exporter must have an establishment Foreign Exporter - Exports or offers for export to the United States (U.S.), a device manufactured or address outside the U.S. Manufacturer - Makes by chemical, physical, biological or other procedures, any article that meets the definition of "device" in Section 201(h) of the Federal Food, Drug, and Cosmetic (FD&C) Act.

Remanufacturer - Processes, conditions, renovates, restores, or does any other act to a finished device that significantly changes the finished device's performance or safety specifications, or in any way changes the intended use.

another Repackager - Packages finished devices from bulk or repackages devices made by manufacturer into different containers (excluding shipping containers). के क Relabeler - Changes the content of the labeling from that supplied from the original manufacturer distribution under the establishment's own name. A relabeler does not include establishments that not change the original labeling but merely add their own name.

Reprocessor of Single Use Devices - Performs remanufacturing operations on a single use device.

Specification Developer - Develops specifications for a device that is distributed under the establishment's own name but performs no manufacturing operations on the device. This includes establishments that in addition to developing specifications also arrange for the manufacturing of devices labeled with another establishment's name by a contract manufacturer. U.S. Manufacturer of Export Only Devices - Manufactures medical devices that are not sold in the U.S. and are manufactured solely for export to foreign countries.

Item 7. Signature of Official Correspondent - The signature of the designated official correspondent.

corres-Item 8. Typed/Printed Name and Title - Type or print the name and title of the designated official pondent.

General Address Instructions:

Enter the number and street - DO NOT use postal box or rural route numbers.

City - Enter the city in which the owner/operator or establishment is located.

State - Enter the two-character state code of the U.S. Postal Service for the State, territory, or possession.

prefecture, region, territory) names in which ZIP Code +4 - Enter the U.S. postal ZIP Code +4 (if known). Foreign State - Enter the foreign state (i.e., province, pr

establishment is located.

Postal Code - Enter the foreign country postal code.

Foreign Country - Enter the full foreign country name

Public reporting burden for this collection of information is estimated to average .25 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to:

| An agency may not conduct or sponsor, and a person is not An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Food and Drug Administration Center for Devices and Radiological Health (HFZ-308) 9200 Corporate Blvd. Rockville, MD 20850-4015

FORM FDA 2892 (6/05)

Worksheet for Imported Footwear

1.	Manufacturer's style number:	A.	Percent of external surface area of upper	(including leather
2.	Importer's style and/or stock number:		reinforcements and accessories) which is	leather
3.	Percentage of external surface area of upper (excluding reinforce-			%
	ments and accessories) which is:	B.	Percent of external surface area of up	per (including all
	Leather a %		reinforcements and accessories) which	
	Composition leather b%		plastics:	
	Rubber and/or plastics c%		2	%
	Textile materials d%	· C.	Percent by weight of rubber and/or plast	ics is:
	Wood e%	-	referred weight of robbet and or pass	96
		D	Percent by weight of textile material(s) p	lus rubber and/or
		D.	A SOLUTION STATE S	as rabber androi
	Other (Specify:) g%		plastics is:	10
	TOTAL 100 %	_		%
4.	Percentage of external surface area of outersole (excluding rein-	Ε.	Is it waterproof?	Yes No
	forcements and accessories) which is:	F.	Does it have a protective metal toe cap?	
	Leather a%		Will it cover the wearer's ankle bone?	Yes No
	Composition leather b %	H.	Will it cover the wearer's knee cap?	☐ Yes ☐ No
	Rubber and/or plastics c %	1.	(RESERVED)	
	Textile materials d%	J.	Is it designed to protect against water, oil,	grease, chemicals,
	Wood e%		or cold or inclement weather?	☐ Yes ☐ No
	Furskin f%	K.	Is it a slip-on?	☐ Yes ☐ No
	Other (Specify:) g%	L	Is it a downhill or cross country skiboot?	☐ Yes ☐ No
	TOTAL 100 %	M.	Is it serious sports footwear other than skit	boots? (See Defini-
5.	In answering 3 or 4, did you consider any pieces to be "accesso-		tion Sheet)	☐ Yes ☐ No
	ries" or "reinforcements"?	N.	Is it a tennis, basketball, gym, or training	shoe or the like?
				☐ Yes ☐ No
	—If yes, which ones?	Ο.	Is it made on a base or platform of wood	7
				☐ Yes ☐ No
		P.	Does it have open toes or open heels?	
			Is it made by the (lipped insole) welt cons	
6.	Regarding the leather pieces (if any) included in 3(a) or 4(a) above,	-	эт постору от рургог этогу, того	☐ Yes ☐ No
	are they coated or laminated with rubber and/or plastics?	R.	Is there a leather outersole attached to I	111-111
	□ Coated □ Laminated □ Neither □ Not Applicable	15.		Yes No
	If coated, is the coating less than 0.1 mm thick?	S.	Is it wom exclusively by men, boys or you	
	☐ Yes ☐ No	٥.	* * * * * * * * * * * * * * * * * * * *	☐ Yes ☐ No
	If no, how thick is the coating to the nearest hundredth of a	т.		ADDRESS CONTRACTOR AND
	millimeter?	9.0	Is it made by an exclusively adhesive cons	
	0 mm			Yes No
7.	You may skip parts I, II and III of this section if you choose to answer	U.		i, by weight, are
	all questions A through Z.		vegetable fibers:	%
	I. If the upper is leather (i.e., 3a accounted for the largest			(Company (Comp
	percent in your response to question 3), then answer ques-	V.	Is it disposable, i.e., intended for one-time	
	tions F. G. L. M. O. Q. R. S and X.			☐ Yes ☐ No
				Yes No
			, , , ,	☐ Yes ☐ No
	the largest percent in your response to question 3), then:	Y.	Are the sole and upper made of wool felt	
	i. if the outer sole is rubber or plastics, answer questions B,			☐ Yes ☐ No
	E, F, G, H, J, K, L, M, N, O, P and W.	Ζ.		
	ii. if the outer sole is not rubber or plastic, STOP.		upper?	Yes No
	III. If the upper is textile materials (i.e., 3d accounted for the			
	largest percent in your response to question 3), then:		lete Items 8-10 if classified under any	
	i. if the outer sole is rubber or plastics, answer questions A.	200000	1.99.80 6402.19.05 6404.19.40	6404.19.60
	C, J, K, M, N, P and T.		2.99.18 6402.19.15 6402.30.30	6402.91.40
	ii. if the outer sole is leather or composition leather, answer	040	4.19.35 6402.99.30 6404.11.40	6404.11.60
	questions C and D.	8. Doe	es the shoe have a foxing or foxing-like ban	d?
	iii. if the outer cole is textile materials, answer questions U,	ф. DO		Yes No
	Y and Z.	0 Dar	es the sole overlap the upper other than just	
	If the item does not fully meet any of the foregoing conditions,			
	answer all of the following questions (A through Z)			Yes No
			e foxing or foxing-like band (if any) almost w	
		olas	tics7	☐ Yes ☐ No

U.S. DEPARTMENT OF HOMELAND SECURITY Bureau of Customs and Border Protection

OMB No. 1651-0098 See back of form for Paperwork Reduction Act Notice.

NORTH AMERICAN FREE TRADE AGREEMENT CERTIFICATE OF ORIGIN

Please print or type	19	CFR 181.11	181 22				
1. EXPORTER NAME AND ADDRESS	13	2. BLANKET					
		FROM					••••••
		T KOW					
TAX IDENTIFICATION NUMBER:		TO		••••••		***************************************	
3. PRODUCER NAME AND ADDRESS		4. IMPORTER	R NAME AND ADDRE	SS			
TAX IDENTIFICATION NUMBER:		TAX IDENTI	FICATION NUMBER:				
5. DESCRIPTION OF GOOD(S)			6. HS TARIFF CLASSIFICATION NUMBER	7. PREFERENCE CRITERION	8. PRODUCER	9. NET COST	10. COUNTR OF ORIGI
		M:					
I CERTIFY THAT: THE INFORMATION ON THIS DOCUMENT IS TRUE REPRESENTATIONS. I UNDERSTAND THAT I AM CONNECTION WITH THIS DOCUMENT;	LIABLE FOR ANY F	ALSE STATE	MENTS OR MATE	RIAL OMISS	SIONS MADE	ON OR IN	
 I AGREE TO MAINTAIN AND PRESENT UPON RE INFORM, IN WRITING, ALL PERSONS TO WHOM T OR VALIDITY OF THIS CERTIFICATE; 	HE CERTIFICATE \	WAS GIVEN C	F ANY CHANGES	THAT COU	LD AFFECT	THE ACCURAC	Υ
THE GOODS ORIGINATED IN THE TERRITORY OF SPECIFIED FOR THOSE GOODS IN THE NORTH A 411 OR ANNEX 401, THERE HAS BEEN NO FURTH PARTIES: AND	MERICAN FREE TE	RADE AGREE	MENT AND UNIE	SS SPECIEI	CALLY EYEN	DTED IN ADTI	CLE
PARTIES; AND THIS CERTIFICATE CONSISTS OF	PAGES, INCLUD	ING ALL ATT	ACHMENTS.				
11a. AUTHORIZED SIGNATURE		11b. COMPAN					
44 NAME (8)							
11c. NAME (Print or Type) 11.		11d. TITLE					
11e. DATE (MM/DD/YYYY)	11f. TELEPHONE NUMBER	(Voice)		(Fac	simile)		

PAPERWORK REDUCTION ACT NOTICE: This information is needed to carry out the terms of the North American Free Trade Agreement (NAFTA). NAFTA requires that, upon request, an importer must provide CBP with proof of the exporters written certification of the origin of the goods. The certification is essential to substantiate compliance with the rules of origin under the Agreement. You are required to give us this information to obtain a benefit.

The estimated average burden associated with this collection of information is 15 minutes per respondent or recordkeeper depending on individual circumstances. Comments concerning the accuracy of this burden estimate and suggestions for reducing this burden should be directed to Bureau of Customs and Border Protection, Information Services Branch, Washington, DC 20229, and to the Office of Management and Budget, Paperwork Reduction Project (1651-0098), Washington DC 20503.

NORTH AMERICAN FREE TRADE AGREEMENT CERTIFICATE OF ORIGIN INSTRUCTIONS

For purposes of obtaining preferential tariff treatment, this document must be completed legibly and in full by the exporter and be in the possession of the importer at the time the declaration is made. This document may also be completed voluntarily by the producer for use by the exporter. Please print or type:

- State the full legal name, address (including country) and legal tax identification number of the exporter. Legal taxation number is: in Canada, employer number or importer/exporter number assigned by Revenue Canada; in Mexico, federal taxpayer's registry number (RFC); and in the United States, employer's identification number or Social Security Number.
- Complete field if the Certificate covers multiple shipments of identical goods as described in Field #5 that are imported into a NAFTA country for a specified period of up to one year (the blanket period). "FROM" is the date upon which Certificate becomes applicable to the good covered by the blanket Certificate (it may be prior to the date of signing this Certificate). "TO" is the date upon which the blanket period expires. The importation of a good for which preferential treatment is claimed based on this Certificate must occur FIELD 2: between these dates.
- State the full legal name, address (including country) and legal tax identification number, as defined in Field #1, of the producer. If more than one producer's good is included on the Certificate, attach a list of additional producers, including the legal name, address (including country) and legal tax identification number, cross-referenced to the good described in Field #5. If you wish this information to be confidential, it is acceptable to state "Available to CBP upon request". If the producer and the exporter are the same, complete field with "SAME". If the producer is unknown, it is acceptable to state "UNKNOWN". FIFI D 3:
- State the full legal name, address (including country) and legal tax identification number, as defined in Field #1, of the importer. If the importer is not known, state "UNKNOWN"; if multiple importers, state "VARIOUS". FIELD 4:
- Provide a full description of each good. The description should be sufficient to relate it to the invoice description and to the Harmonized System (H.S.) description of the good. If the Certificate covers a single shipment of a good, include the invoice number as shown on the commercial invoice. If not known, indicate another unique reference number, such as the shipping order number. For each good described in Field #5, identify the H.S. tariff classification to six digits. If the good is subject to a specific rule of origin in Annex 401 that requires eight digits, identify to eight digits, using the H.S. tariff classification of the country into whose territory the good is FIELD 5:
- FIELD 6: imported.
- For each good described in Field #5, state which criterion (A through F) is applicable. The rules of origin are contained in Chapter Four and Annex 401. Additional rules are described in Annex 703.2 (certain agricultural goods), Annex 300-B, Appendix 6 (certain textile goods) and Annex 308.1 (certain automatic data processing goods and their parts). NOTE: In order to be entitled to preferential tariff treatment, each good must meet at least one of the criteria below. FIELD 7:

Preference Criteria

- The good is "wholly obtained or produced entirely" in the territory of one or more of the NAFTA countries as referenced in Article 415. Note: The purchase of a good in the territory does not necessarily render it "wholly obtained or produced". If the good is an agricultural good, see also criterion F and Annex 703.2. (Reference: Article 401(a) and 415)
- The good is produced entirely in the territory of one or more of the NAFTA countries and satisfies the specific rule of origin, set out in Annex 401, that applies to its tariff classification. The rule may include a tariff classification change, regional value-content requirement, or a combination thereof. The good must also satisfy all other applicable requirements of Chapter Four. If the good is an agricultural good, see also criterion F and Annex 703.2. (Reference: Article 401(b))
- The good is produced entirely in the territory of one or more of the NAFTA countries exclusively from originating materials. Under this criterion, one or more of the materials may not fall within the definition of "wholly produced or obtained", as set out in article 415. All materials used in the production of the good must qualify as "originating" by meeting the rules of Article 401(a) through (d). If the good is an agricultural good, see also criterion F and Annex 703.2. Reference: Article 401(c).
- Goods are produced in the territory of one or more of the NAFTA countries but do not meet the applicable rule of origin, set out in Annex 401, because certain non-originating materials do not undergo the required change in tariff classification. The goods do nonetheless meet the regional value-content requirement specified in Article 401(d). This criterion is limited to the following two circumstances:
 - The good was imported into the territory of a NAFTA country in an unassembled or disassembled form but was classified as an assembled good, pursuant to H.S. General Rule of Interpretation 2(a), or
 - The good incorporated one or more non-originating materials, provided for as parts under the H.S., which could not undergo a change in tariff classification because the heading provided for both the good and its parts and was not further subdivided into subheadings, or the subheading provided for both the good and its parts and was not further subdivided.

NOTE: This criterion does not apply to Chapters 61 through 63 of H.S. (Reference: Article 401(d))

- Certain automatic data processing goods and their parts, specified in Annex 308.1, that do not originate in the territory are considered originating upon importation into the territory of a NAFTA country from the territory of another NAFTA country when the most-favored-nation tariff rate of the good conforms to the rate established in Annex 308.1 and is common to all NAFTA countries. (Reference: Annex 308.1)
- The good is an originating agricultural good under preference criterion A, B, or C above and is not subject to a quantitative restriction in the importing NAFTA country because it is a "qualifying good" as defined in Annex 703.2, Section A or B (please specify). A good listed in Appendix 703.2B.7 is also exempt from quantitative restrictions and is eligible for NAFTA preferential tariff treatment if it meets the definition of "qualifying good" in Section A of Annex 703.2. NOTE 1: This criterion does not apply to goods that wholly originate in Canada or the United States and are imported into either country. NOTE 2: A tariff rate quota is not a quantitative restriction.
- For each good described in Field #5, state "YES" if you are the producer of the good. If you are not the producer of the good, state "NO" followed by (1), (2), or (3), depending on whether this certificate was based upon: (1) your knowledge of whether the good qualifies as an originating good; (2) your reliance on the producer's written representation (other than a Certificate of Origin) that the good qualifies as an originating good; or (3) a completed and signed Certificate for the good, voluntarily provided to the exporter by the producer. FIELD 8:
- For each good described in field #5, where the good is subject to a regional value content (RVC) requirement, indicate "NC" if the RVC is calculated according to the net cost method; otherwise, indicate "NO". If the RVC is calculated over a period of time, further identify the beginning and ending dates (MM/DD/YYYY) of that period. (Reference: Article 402.1, 402.5). FIFLD 9
- Identify the name of the country ("MX" or "US" for agricultural and textile goods exported to Canada; "US" or "CA" for all goods exported to Mexico; or "CA" or "MX" for all goods exported to the United States) to which the preferential rate of CBP duty applies, as set out in Annex 302.2, in accordance with the Marking Rules or in each party's schedule of tariff elimination. FIELD 10:
 - For all other originating goods exported to Canada, indicate appropriately "MX" or "US" if the goods originate in that NAFTA country, within the meaning of the NAFTA Rules of Origin Regulations, and any subsequent processing in the other NAFTA country does not increase the transaction value of the goods by more than seven percent; otherwise indicate "JNT" for joint production. (Reference: Annex 302.2)
- FIELD 11: This field must be completed, signed, and dated by the exporter. When the Certificate is completed by the producer for use by the exporter, it must be completed, signed, and dated by the producer. The date must be the date the Certificate was completed and signed.

CBP Form 434 (04/97)(Back)

COMPOSITION BREAKDOWN FOR POLYRESIN ITEMS

<u>ITEN</u> <u>SKU</u> ITE			
1. Does the Yes (e item consist of a crushed or ground) No () (If yes, please go to poin	*	e material?
	item consist of a synthetic chemical No () (If yes, stop here. No form		
3. Please in	ndicate the precise geological name	of any stone present in the product	
	aterial used is calcite (eg, Calcium C or ground stone or from a synthetic	· •	the calcite is derived from
5. Is the ste Yes (one or mineral material uniformly bl	ended with the plastic material thro	oughout the article?
6. Please in	ndicate the percentage by weight and	I the percentage by value of each n	naterial within the product.
MATER	IAL	% BY WEIGHT	% BY VALUE

Textile Backpacking Tent Form

To qualify a tent as "sport equipment" under HTSUS# 6306.22.1000, the following criteria must be met. Please describe your backpacking tent by filling in the required information below.

(1) It must be specially designed for the sport of backpacking. () Yes or () No
(2) It must be composed of () nylon, () polyester, () any other fabric of man-made fibers, please specify
(3) If designed for 1 or 2 persons, the tent must meet the following criteria:
(a) Have a floor area of (square feet) that is 45 square feet or less, and
(b) Weigh (pounds) that is 8 1/2 pounds or less, including tent bag and all accessories necessary to pitch the tent, and
(c) Have a carry size of(length x diameter inch) that is 30 inches or less in length and 9 inches or less in diameter. If other than cylindrical in shape, the tent package must not exceed 1,900 cubic inches.
(4) If designed for 3 or 4 persons, the tent must meet the following criteria:
(a) Have a floor area of (square feet) that is 65 square feet or less; and
(b) Weigh (pounds) that is 12 pounds or less, including tent bag and all accessories necessary to pitch the tent and
(c) Have a carry size of(length x diameter inch) that is 30 inches or less in length and 10 inches or less in diameter. If other than cylindrical in shape, the tent package must not exceed 2,350 cubic inches.

Any tent with a floor of more than 65 square feet and a standing height of more than 60 inches is a tent designed for general recreational use.

In any part of the tent package (including tent skin, frame, tent bag, accessories or packing) is not imported in the same shipment with all remaining parts, in order to receive the benefit of the sport equipment classification for such part, the importer must supply Customs with a sample of the complete tent package so a determination of compliance can be made.

ISFWS Form 3-177 Revised 12/03)		U.S. FISH AND WI	LDLIFE SERVICE		Page	of
.M.B. No. 1018-0012 expiration Date: 12/31/200 I. Date of Import/E / 2. I/E License Num 3. Indicate One:	xport: (mm/dd/yyyy)	DECLARATION FOOR EXPORTED FISH OR VERNING TO THE PROPERTY OF T	FATION OF VILDLIFE	Master: House: 9. Transport 10. Bonded	Bill or Bill of Lading	State dion:
13. (indicate one) U.S. Importer U.S. Exporte	r of Record r r er, Shipping Agent or Freight	address / phone number) t Forwarder:	14a. Foreign Suppl (complete nan	ier / Receiver; ne / address / phone		4b
Species Code Official Use)	16a. Scientific Name	175	Foreign CITES Permit Number U.S. CITES Permit Number	18a. Description Code	19a. Quantity / Unit 19b. Total Monetary Value	20. Country of Origin of Animal
Knowingly making a f of Fish or Wildlife ma 1001 and 16 U.S.C. 3 Action/Comments:	` '	or Importation or Exportation enalty provided by 18 U.S.C	21. I certify under pend correct:	Signature Type or Print Name	nformation furnished is	s true and Date



BIG LOTS STORES

Department 30905A 300 Phillipi Rd P.O.Box 28512 Columbus, OH 43228-0512 U.S.A.

Phone: (614)278-3748 Fax: (614) 278-7161

TSCA STATEMENT

Entry# : Invoice#:
Please select the TSCA statement that is applicable for item#:
TSCA + positive certification for shipments subject to TSCA. I certify that the chemical substance on this invoice line complies with all applicable rules or orders under TSCA and am not offering a chemical substance for entry in violation of TSCA or any applicable rule or order there under.
☐ TSCA – negative certification for shipments not subject to TSCA. I certify that the chemical substance on this invoice line is not subject to TSCA.
Signature:
BIG LOTS STORES INC.

VERIFICATION OF UNDERWRITERS LABORATORIES INC. CERTIFICATION (UL / ETL)

I,	(S	igner's name),		(Signer's title) of
Certificates are v	valid for product b	eing sold to Big Lots	that the enclosed Underwriters less Stores, Inc. and/or direct and in and covers the following item(s	ndirect subsidiaries under Purchase Order(s
<u>Item No.</u>		_	<u>le UL file numbers)</u>	
Sworn to this	day of	, 20		
Signature				
Title				
Company				

*ATTACH UL/ETL CERTIFICATES

SOLD BY: (this must match purchase order) Full name and address, fax and phone

COMMERCIAL INVOICE

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INVOICE NO: SHIPPING TERMS:

L/C NO:

PAYMENT TERMS:

CARRIER:

PORT OF DISCHARGE:

SHIP TO

P.O. # Item # and sku # full description of product qty unit price* USD amount

*cost breakdown by piece if it is a set

TOTAL: USD 0.00

SAY US DOLLARS.....

WE HEREBY CERTIFY THAT THE SHIPMENT CONTAINS NO WOOD PACKAGING MATERIALS

DESTINATION:

MANUFACTURERS NAME AND ADDRESS

P.O. # SKU # DEPT # CARTON OF MADE IN XXXXX

> WE CERTIFY THAT INFORMATION SHOWN ABOUT MANUFACTURER IS TRUE AND ACCURATE

FOR AND ON BEHALF OF XXX (Original SIGNATURE)

Forwarders Cargo Receipt

NYK Logistics - NYK Logistics	NYK	Logistics	NYK Logistics -	NYK Logistics
				THE LOGISTICS
401	RWARDER'S CARGO) RECEIPT No.		
Maker/Supplier:			Maker/Supplier	's INVOICE No.
Buyer/Consignee :				
Shipment From:	То:			
Marks & Nos.	Nos. of P'kgs	Supplier's description of goods	Measurement (cbm.)	Weight (kgs.)
				Weight (kgs.)
RECEIVED by NYK Legistics (Heng Kong) I Company) in apparent good order and condition the goods, or the container(s) or package(s) said foreinafter called the Goods) for transportation	, unless otherwise indicates to contain goods herein an	I herein, sentioned		Control of the Contro
the owner of the Goods. The Company is author carries involved in the execution of the transport conditions of the carriers. While the Goods are is Company shall hold the same as a Freight Forw not be responsible for any loss, damage and for may be caused by the willful or grossly negligen agents or employees. In winness whereof, the un Logistics (Hong Kong) Limited has signed the Receipt(s) stated below, all of this tenor and date No., of ORIGINAL FORWARDER'S C	rized to enter into contracts tation subject to all terms a in the custody of the Comp arder and not as a carrier at cleay of the Goods, excer at the Company, its or dersigned, on behalf of N' number of Forwarder's Ca c.	with nd d nd nd nd NYK L such as fifters, VK	(Place and date of issue.) ogistics (Hong Kong) Lin	nited

CERTIFICATE OF LIABILITY INSURANCE

Date (MM/DD/YY):

PRODUCER

AIG General Insurance Company China Limited Shanghai Branch

15F, Bank of China Tower, 200 Yincheng Road Central, Pudong, Shanghai 200120 P.R.CHINA THIS CERTIFICATE IS ISSUED AS A MATTER OF INFORMATION ONLY AND CONFERS NO RIGHTS UPON THE CERTIFICATE HOLDER. THIS CERTIFICATE DOES NOT AMEND, EXTEND OR ALTER THE COVERAGE AFFORDED BY THE POLICIES BELOW.

COMPANIES AFFORDING COVERAGE

COMPANY

AIG GENERAL INSURANCE COMPANY CHINA LIMITED SHANGHAI BRANCH

INSURED

COVERAGES

THIS IS TO CERTIFY THAT THE POLICIES OF INSURANCE LISTED BELOW HAVE BEEN ISSUED TO THE INSURED NAMED ABOVE FOR THE POLICY PERIOD INDICATED, NOTWITHSTANDING ANY REQUIREMENT, TERM OR CONDITION OF ANY CONTRACT OR OTHER DOCUMENT WITH RESPECT TO WHICH THIS CERTIFICATE MAY BE ISSUED OR MAY PERTAIN, THE INSURANCE AFFORDED BY THE POLICIES DESCRIBED HEREIN IS SUBJECT TO ALL THE TERMS, EXCLUSIONS AND CONDITIONS OF SUCH POLICIES. LIMITS SHOWN MAY HAVE BEEN REDUCED BY PAID CLAIMS.

	IIIONO OI DOON IOHIOIHD.	DIMITE CHOWN	MI TIMVE DEELN	REDUCED DI F	AID CLAIMS.	
CO LTR	TYPE OF INSURANCE	POLICY NUMBER	POLICY EFFECTIVE DATE (MM/DD/YY)	POLICY EXPIRATION DATE (MM/DD/YY)	LIMITS	
	GENERAL LIABILITY		MM/DD/YY	MM/DD/YY	GENERAL AGGREGATE	N/A
A	☐ COMMERCIAL GENERAL LIAB.		(AT 12:00 A.M. STANDARD TIME AT	(AT 00:00 A.M. STANDARD TIME	PRODUCTS - COMP/OP AGG	US\$1,000,000
	☑ PRODUCTS LIABILITY	1/1	THE INSURED'S ADDRESS SHOWN ABOVE)	AT THE INSURED'S ADDRESS SHOWN ABOVE)	PERSONAL & ADV INJURY	N/A
	☐ CLAIM MADE ☐ OCCUR	α I α r	1, I V		EACH OCCURRENCE	US\$1,000,000.
	☐ OWNER'S & CONTRACTOR'S PROT	YIVII '	# de 1		FIRE DAMAGE (Any one fire)	N/A
	☐ BROAD FORM VENDORS COVERAGE	7 Trian			MED E PAY (Any one person)	N/A
	AUTOMOBILE LIABILITY ANY AUTO				COMBINED SINGLE LIMIT	\$
	☐ ALL OWNED AUTOS				BODILY INJURY (Per person)	\$
	☐ SCHEDULED AUTOS ☐ HIRED AUTOS				BODILY INJURY (Per accident)	\$
	□ NON-OWNED AUTO				PROPERTY DAMAGE	\$
	GARAGE LIABILITY				AUTO ONLY - EA ACCIDENT	
	☐ ANY AUTO				OTHER THAN AUTO ONLY:	
	□				EACH ACCIDENT	\$
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	EXCESS LIABILITY				EACH OCCURRENCE	\$
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	WORKERS COMPENSATION AND				WC STATU- OTH-	
	EMPLOYERS' LIABILITY				TORY LIMITS ER	
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	THE PROPRIETOR/ PARTNERS/EXECUTIVE □INCL				EL DISEASE - POLICY LIMIT	\$
	OFFICERS ARE: EXCL				EL DISEASE – EA EMPLOYEE	\$
	OTHER					

Description of Operations/Locations/Vehicles/Special Items

This Insurance is primary with respect of Products Covered in this Policy and subject to the Policy

terms and conditions

Jurisdiction/Territory: Worldwide

SPECIAL CONDITIONS: DISCONTINUED SUPPLIER'S EXTENSION-3 YEARS; &

DEFENSES COSTS IN ADDITION TO LIMIT OF LIABILITY

CERTIFICATE HOLDER

Additional Insured:

Big Lots, Inc. and its direct and indirect subsidiaries

300 Phillipi road, Columbus, Ohio 43228, USA

CANCELLATION

SHOULD ANY OF THE ABOVE DESCRIBED POLICIES BE CANCELLED BEFORE THE EXPIRATION DATE THEREOF, THE ISSUING COMPANY WILL ENDEAVOR TO MAIL 30 DAYS WRITTEN NOTICE TO THE CERTIFICATE HOLDER NAMED TO THE LEFT, BUT FAILURE TO MAIL SUCH NOTICE SHALL IMPOST NO OBLIGATION OR LIABILITY OF ANY KIND UPON THE COMPANY, ITS AGENTS OR REPRESENTATIVE. 10 DAYS WRITTEN NOTICE WILL BE GIVEN FOR NON-PAYMENT OF PREMIUM.

AUTHORIZED REPRESENTATIVE

Certificate Alert: to check on the validity of this certificate or the underlying policy or to receive an e-copy, please email us with the certificate number at china.products@aig.com.

Certificate No.: Certificate Specimen

Single Country Declaration									
or country, insular posses United States, of fabricat the United States and/or	(name), declare that the articles listed relates are wholly the growth, products ssion of the United States, or were asseted components which are in the whole the single foreign territory of country, of information set forth in this declaration and belief.	s, or manufacture of a sing mbled in the single foreig product of or insular of the United St	gle foreign territory in territory or country, or i ates, as identified	insular possession of the					
(Country)									
Marks of identification, numbers	Description of article and Qty (Item #, SKU #, P.O. #)	Country of origin	Date of Export						

Signature:	Company:	
Name:	Address:	
Гitle:		
Date:	Invoice #:	

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	Phone/Fax			Phone/Fax				moma use only		Weight (lbs) Gaude (metric)			
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	Contact Name			Contact Name						Price Quoted notil (Order Date)	(Order Date)		
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Trademark Sublicense Agreement Form

Big Lots uses original brands on packaging, including Shop Basics, ProSource, Village Green, BBQ Grates, and Classic Quarters (see complete list below). Vendors are required to complete and return a Trademark Sublicense Agreement form (pages 43-45) for items using these brands on packaging. The Agreement gives permission to the vendor to use the brand, and explains vendor use limitations.

Big Lots Brands

BBQ GRATES BIG LOTS PRIVATE LABEL BRIDGEPORT (Office & Juvenille) CAMPLIFE CLASSIC QUARTERS CLIMATE KEEPER COMFEES COUNTER COOK DAKIN **E SOURCE** FRESH FINDS IT'S A KEEPER **GREAT GATHERINGS** LIVING COLORS ONCE UPON A TIME PRO SOURCE PRO SOURCE BASICS QUAL STAR READY SET ROOM SHOP BASICS SOUNDBODY STYLE ELEMENTS VILLAGE GREEN WESTMINSTER CLASSICS WILSON & FISHER

TRADEMARK SUBLICENSE AGREEMENT

This Trademark Sublicense Agreement ("Agreement") contains a sublicense from Big Lots Stores, Inc., an Ohio corporation having its principal place of business at 300 Phillipi Road, Columbus, Ohio 43228 ("Big Lots"), for use of trademarks specified herein ("Trademarks") and will only become effective upon your assent to all terms herein, as evidenced by your execution of this Agreement. If you do not accept all terms of this Agreement, you shall not be permitted to use the Trademarks for any purpose or have any other right in the Trademarks or goods bearing the Trademarks. You, the company or organization set forth below, are referred to as the "Sublicensee" in this Agreement.

1. SUBLICENSE.

(a) Sublicense Grant. Upon the acceptance of this Agreement, Big Lots grants to Sublicensee, and Sublicensee accepts, a nonexclusive, nontransferable, personal right to use, under the express terms of this Agreement, only the following Trademarks and only for the purposes of labeling goods ordered by Big Lots with a valid purchase order placed by Big Lots for the following goods:

Trademark	DESCRIPTION OF SPECIFIC GOODS	ITEM No.

(b) Additional Sublicensee Restrictions. Nothing in this Agreement shall give Sublicensee any right, title or interest in the Trademarks, including trade dress, copyrights, package design or any other property right, (or any other trademark or right of Big Lots or its affiliated companies), other than the sublicense rights expressly granted in Subsection 1(a) above. Sublicensee shall not assign, transfer or sublicensee this Agreement (or any right granted herein) without the prior written consent of Big Lots. Sublicensee agrees not to use the Trademarks in any way that would disparage or injure Big Lots' reputation for high quality. ADDITIONALLY, LICENSEE AGREES THAT GOODS MANUFACTURED AND/OR LABELED BY SUBLICENSEE SHALL NOT BE DEEMED AUTHORIZED OR GENUINE GOODS UNTIL: (1) BIG LOTS HAS EXPRESSLY DECLARED SUCH GOODS TO BE GENUINE IN WRITING, OR (2)(i) BIG LOTS HAS RECEIVED SUCH GOODS AT ITS DISTRIBUTION CENTER, (ii) HAS INSPECTED THE GOODS TO CERTIFY THEIR QUALITY AND (iii) THE GOODS ARE NOT MATERIALLY DIFFERENT IN ANY MANNER FROM GOODS EXPRESSLY AUTHORIZED BY BIG LOTS PREVIOUSLY.

 DISTRIBUTION LABEL. If applicable, for the mutual benefit of the parties, Big Lots hereby grants Sublicensee permission to print the following text on the packaging of goods to be sold only in Big Lots' stores in connection with the Trademark(s) specified below:

> Distributed By: Big Lots Stores, Inc.; P.O. Box 28523; Columbus, OH 43228-0523 V#000000 MADE IN CHINA ITEM #000000

Trademark	DESCRIPTION OF SPECIFIC GOODS	ITEM No.

Trademark Sublicense Agreement - 0609200806092008

3. ADDITIONAL USE RESTRICTIONS & GUIDELINES.

- (a) The Trademarks may be used only to identify Big Lots as the source of Big Lots goods, as specifically provided in Subsection 1(a) above.
- (b) Sublicensee may use the Trademarks so long as such use complies with this Agreement, including:
 - Sublicensee shall only use the Trademarks on goods sold or provided directly to Big Lots and not in any other manner;
 - (ii) The Trademarks must appear exactly as provided by Big Lots' direction and not in any other manner;
 - (iii) The Trademarks shall not be altered in any way, including but not limited to, skewing, changing the color, rotating, separating logo elements or changing a typeface;
 - (iv) Sublicensee may reproduce the Trademarks only as expressly permitted by this Agreement;
 - (v) Sublicensee may not display the Trademarks in any manner that implies affiliation with, sponsorship, or endorsement by any person, company or organization other than Big Lots;
 - (vii)Sublicensee may not display the Trademarks in any manner that disparages Big Lots or its goods and services, infringes Big Lots' intellectual property, or violates any state, federal, or international laws; and
 - (vii) Sublicensee shall adhere to all quality control standards imposed by Big Lots.
- (c) Protecting the Trademarks. Sublicensee acknowledges Big Lots' rights in and/or ownership of (and/or its affiliated companies' rights in and/or ownership of) the Trademarks. Sublicensee shall not use the Trademarks in a manner that does derogates from Big Lots' and its affiliated companies' rights in the Trademarks and will take no action that will interfere with or diminish Big Lots' and its affiliated companies' rights in the Trademarks. All uses of the Trademarks by Sublicensee will inure to the benefit of Big Lots and its affiliated companies. Sublicensee will not use the Trademarks in any way as an endorsement or sponsorship by Big Lots of a third party or Sublicensee product, website, service, information or other content.

4. QUALITY STANDARD; INSPECTION AND APPROVAL.

- (a) Standard. Sublicensee agrees to maintain the quality of Sublicensee's use of the Trademarks that meets or exceeds industry standards.
- (b) Inspection. Upon reasonable request from Big Lots, Sublicensee shall notify Big Lots of all of Sublicensee's use of the Trademarks and furnish Big Lots with suitable specimens of Sublicensee's use of the Trademarks. Big Lots may review Sublicensee's use of the Trademarks periodically to evaluate Sublicensee's compliance with the quality standards provided by Big Lots and as described in this Agreement. Sublicensee shall immediately remedy any deficiencies in its use of the Trademarks, upon discovery of the same or upon notice from Big Lots and destroy all goods and packaging bearing the Trademarks in contradiction to this Agreement.
- (c) Sublicensee's Liability. Sublicensee shall fully indemnify, defend (at Big Lots' option) and hold harmless Big Lots from and against any claims relating to Sublicensee's use of the Trademarks that does not comply with this Agreement or any state, federal or international law or regulation.
- 5. LIMITATION OF LIABILITY. BIG LOTS MAKES NO WARRANTIES OF ANY KIND, EITHER EXPRESS OR IMPLIED, WITH RESPECT TO THE TRADEMARKS OR THE RELATED GOODS. IN NO EVENT SHALL BIG LOTS BE LIABLE FOR ANY CONSEQUENTIAL, INCIDENTAL, OR SPECIAL DAMAGES (INCLUDING LOSS OF BUSINESS PROFITS) ARISING FROM OR RELATED TO SUBLICENSEE'S USE OF THE TRADEMARKS, EVEN IF BIG LOTS HAS BEEN ADVISED OF THE POSSIBILITY OF SUCH DAMAGES. IN ALL CASES, BIG LOTS' AGGREGATE LIABILITY SHALL BE LIMITED TO DIRECT DAMAGES NOT TO EXCEED US\$1,000.00.

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- 6. TERM AND TERMINATION. The date of acceptance of this Agreement by Sublicensee shall be the Effective Date of this Agreement. The term of this Agreement shall be for a period of one (1) month from the Effective Date and will automatically renew for successive one (1) month terms until either party gives the other party thirty (30) days prior written notice of termination. Big Lots shall have the right to immediately terminate this Agreement with cause if it determines, in its sole discretion, that Sublicensee is not using the Trademarks in compliance with this Agreement. Any violation of this Agreement by Sublicensee or other misuse of the Trademarks or Big Lots' (or its affiliated companies') other intellectual property rights shall automatically and immediately terminate this Agreement, with or without notice from Big Lots. From and after termination or expiration of this Agreement, Sublicensee shall cease and desist from all use of the Trademarks and shall immediately destroy all goods and packaging bearing the Trademarks.
- 7. NOTICES. All notices under this Agreement shall be in writing and shall be deemed given if sent by:
 - (a) confirmed facsimile to Sublicensee at the facsimile number set forth below,
 - (b) overnight courier, with confirmation of receipt, to Big Lots Stores, Inc., at 300 Phillipi Road, Columbus, Ohio, USA 43228, Attention: Package Design Department, with a copy to Big Lots' General Counsel at the same address.
- 8. MISCELLANEOUS. The parties agree that, in lieu of any royalty for use of the Trademarks, Sublicensee will factor the value of its use of the Trademarks in connection with this Agreement into the purchase price of the related goods being charged to Big Lots. This Agreement contains the entire agreement of the parties with respect to the subject matter hereof and shall not be amended except by a written agreement subsequent to the Effective Date and signed by authorized representatives of each party. This Agreement shall not modify any purchase order placed by Big Lots to Sublicensee or its affiliates, but the terms of this Agreement shall prevail in the event, and only to the extent, that they conflict with the express terms of a Big Lots purchase order. This Agreement shall be governed by and construed in accordance with the laws of the State of Ohio. Sublicensee hereby consents to the exclusive jurisdiction and venue in the state and federal courts sitting in Franklin County, Ohio, USA, and waives all objections to lack of personal jurisdiction and an inconvenient forum. If either party employs attorneys to enforce any rights arising out of or related to this Agreement, the prevailing party shall be entitled to recover its reasonable attorneys' fees, costs, and other expenses. No waiver of any breach of any provision of this Agreement shall constitute a waiver of any prior, concurrent or subsequent breach of the same or any other provision hereof, and no waiver shall be effective unless made in writing and signed by an authorized representative of the waiving party. If any provision of this Agreement shall be held by a court of competent jurisdiction to be illegal, invalid or unenforceable, the remaining provisions shall remain in full force and effect. This Agreement shall not be construed as creating a partnership, joint venture or agency relationship or as granting a franchise. This Agreement is made in Columbus, Ohio, USA, as of the date written below by an officer of the Sublicensee authorized to execute contracts of this nature on behalf of his/her company.

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Trademark Sublicense Agreement - 0609200806092008

Bureau Veritas Locations



Move Forward with Confidence



Bureau Veritas Consumer Products Services

Americas

Canada

Bureau Veritas Consumer Products Services, Inc.9 8220 Bayview Avenue, Suite 207 Thomhill, Ontario L3T 2S2 Canada Tel: 905 771 5722 / Fax: 905 771 5724 Contact: Steven Solarik Email: steven.solarik@ca.bureauveritas.com

Guatemala

Bureau Veritas Consumer Products Services Guatemala S.A.L 5 Av. 5-55 zona 14, Edificio Europlaza Torre IV, Nivel 15 C.P. 01014, Guatemala, Guatemala, Tel: 502 2385 3741 / Fex: 502 2385 3748 Contact: Lillian Peregrina Email: lillian.peregrina@gt.bureauveritas.com

Bureau Veritas Consumer Products Services Mexico, SA DE CV.1-Circuito del Sol #3905-1. Colonia. Nuevo American Puebla, Pue. Mexico CP 72400 Tel: 52 222 266 0561 / Fax: 52 222 266 0563 Contact: Carlos J. Lozano Email: carlos.lozano@mx.bureauventas.com

California

Bureau Veritas Consumer Products Services, Inc.9 425 East Colorado Street, Suite 660 Glendale, California 91205, USA Tel: 818 549 2470 / Fax: 818 549 2471 Contact: Mari Miller Email: mari.miller@us.bureauveritas.com

Bureau Veritas Consumer Products Services, Inc.9 Social Accountability Division 525 Brannan Street, Suite 208 San Francisco, California 94107, USA Tel 415 537 6980 Email: bucps socialaccountability @us.bureauveritas.com

Illinois (consulting office) Bureau Veritas Consumer Products Services, Inc. Butterfield Office Plaza, 2625 W. Butterfield Rd, Ste 313 W Oak Brook, Illinois 60523, USA Tel: 630 203 4400 / Fex: 630 203 4399 Contact: Joan Mattson Email: joan.mattson@us.bureauveritas.com

Massachusetts

Bureau Veritas Consumer Products Services, Inc. 350 Myles Standish Blvd. Taunton, Massachusetts 02780, USA Tel Tall free: 1 800 580 8353 or 508 844 2700 Fex: 508 844 2869 Email: info@us.bureauveritas.com

Curtis-Straus LLC, A Bureau Veritas Company 4 527 Great Road Tel: 978 486 8880 / Fex: 978 486 8828 Email: sales@curtis-straus.com

Bureau Veritas Consumer Products Services, Inc. 4 100 Northpointe Parkway Buffelo, New York 14228, USA Tel: Toll-free 1 800 277-3300 or 716 505 3300; Fax: 716 505 3301 Email: info@us bureauveritas.com

Europe

France

Bureau Veritas Consumer Products Services France 4 Rue John Hadley, BP 20431 59658 Villeneuve d'Ascq, France Tel: 33 3 20 463 456 / Fax: 33 3 20 46 9906 Email: contact bycostrance@fr.bureauveritas.com

LCIE, A Bureau Veritas Company L 33, avenue du General Leclero 92260 Fonteney-aux-Roses, France Tel: 33 1 40 95 60 60 / Fax: 33 1 40 95 86 56 Email: chris.baird@fr.bureauveritas.com

LCIE HAZTEC

Adresse 4, Av. de la Baltique - Bâtiment B 91962 Courtaboeur Tel: +33 1 69 07 87 78 / Fax: +33 1 69 28 81 44 Contact - Stephane Ranchon Email: stephane.ranchon@lcie.fr

Germany
Bureau Veritas Consumer Products Services Germany W. - Hennemann - Str. 8 D-19061 Schwerin, Germani Tel: 49 40 5302084-0 / Fax: 49 40 5302084 80 Contact: Dr. Manfred Mayer Email: cps-schwerin@de.bureauventas.com

Bureau Veritas Consumer Products Services Germany-Georg-Wilhelm Str. 183, D-21107 Hamburg Tel: 49 40 5302084-0 / Fex: 49 40 5302084-19 Contact: Dr. Manfred Mayer Email: cps-hamburg@de.bureauveritas.com

Bureau Veritas E & E Product Services GmbH 4 Business park A96, 86842 Tuerkheim, Germany Tel: +49 (0) 8245 96810-0 / Fax: +49 (0) 8245 96810-99 Email: horst.haug@innova-ps.de

Bureau Veritas Consumer Products Services Spain? Francisca Delgado, 11 Parque Empresarial Arroyo de la Vega 28 108, Alcobendas, Madrid, Spain Tel. (34) 91 270 22 00 / Fax: (34) 91 270 22 98 Email: CPDEspMail@es.bureauveritas.com

Turkey

Bureau Veritas Consumer Products Services Test Laboratuvariari Ltd. Sti.4 Mehmet Akif Mah, Express Cad Can Sok, No. 31, Kat: 3, Ikitelli, Istanbul, Turkey Tel: +90 212 494 35 35 / Fex: +90 212 494 35 60 Contact: Atanur Dikmen Email: atanur.dikmen@bvcps.com.tr

United Kingdom Bureau Veritas Consumer Products Services UK Ltd. 9 2nd Floor, Tower Bridge Court, 224-226 Tower Bridge Rd. London, SEI 2TX, United Kingdom Tel: 44 207 550 8900 / Fax: 44 207 089 7085 Email: bysales@uk.bureauveritas.co

Bureau Veritas Consumer Products Services UK Ltd. 4 Oakhurst House, 57, Ashbourne Road Derby DE22 3FS, United Kingdom Tel: +44 1 332 36 3100 / Fex: +44 1 332 36 3700 Contacts: Alan Kirwilliam Director Sales & Marketing UK Chris Davey, Director Operations UK Email: BVSales@uk.bureaverites.com

Bureau Veritas Consumer Products Services UK Ltd. 4 31 Kingsland Grange, Woolston Warrington, Cheshire WA1 4RW United Kingdom Tel: +44 1925 854350 / Fax: +44 1925 851654 Contacts: Alan Kinwilliam Director Sales & Marketing UK Chris Davey, Director Operations UK Email: BVSales@uk.bureaveritas.com



Asia

Bangladesh

Bureau Veritas Consumer Products Services (BD) Ltd.1-Bashati Horizon, Flat # 8-6, 6th Floor, House #21, Road #17 Banani C/A, Dhaka 1213, Bangladesh. Tel: 88-02-8860342 / Fax: 88-02-9893097 Contact: Dr. Simon Mak Email: simon.mak@bd.bureauveritas.com

China

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Bureau Veritas Consumer Products Services L (softlines/analytical) No.168, Guanghua Road, Zhuanqiao Town, Minhang Shanghai, China. Post Code: 201108 Tel: (86) 21 24081888 / Fax: (86) 21 64890042 Contact: Anita Lui / Judy Yu / Julie Jiang Email: byops_sh_bd@cn.bureauveritas.com

Bureau Veritas Consumer Products Services (Inspection, Audit & Assessment Services office) Room 410, 2nd building, No.351, ZhaoJiaBang Road, Xultui District, Shanghai, China, Post Code: 200032 Tel: (86) 21-6418-1211 / Fax: (86) 21-6418-3853 Email: bvcps.inspection-sh@cn.bureauveritas.com

Advance Data Technology Corporation, i A Bureau Veritas Company 2F, Building C, No. 1618 Yishan Road, Shanghai, China Tel: (86) 21-6465-9091 / Fax: (86) 21-6465-9092 Contact: Wallace Pan Email: wallace_pan@adt.com.tw ADT Website: www.adt.com.tw

LCIE China (F5, Building 10, No.489 North Tibet Road 200071, Shanghai, China Tel: (86) 21-6605-8286 / Fax: (86) 21-5662-1192 Contact: Tarik Mahammed Email: tank.mahammed@fr.bureauventas.com

Bureau Veritas Consumer Products Services L (hardlines) 1st & 4th Floor, B Building, Minlida Industrial Building, Honghualing Industrial Park, Liu Xian Road, Xili, Narshan District, ShenZhen, Guangdong Province, China, Post Code: 518055 Tet (86) 755-8600-0151 Sample pickup: (86) 755-8600-0151 ext. 101 or 102 Fax: (86) 755-8600-0157 Cantact: Wingo Chee E-mail: bvcps_szinfo@cn.bureauveritas.com

Bureau Venitas Consumer Products Services (toys) 4th Floor, Block 1, Dayuan Industrial District South, Liu Xian Da Dao, XiLi, Nanshan District, Shenzhen, Guengdong Province, China, Post Code: 518055 Tel: (86) 755-8602-7802 / Fax: (86) 755-8602-7955 Contact: Casio Chen Email: bvcps_szinfo@cn.bureauveritas.com

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Bureau Ventas Consumer Products Services ((softines) No. 183, Shi Nan Road, Mei Lin Plaza Block B. Dong Chong, Pan Yu, Guangzhou, Guang Dong Province, China Tel: (86) 020-22902088 Ext. 301/320 Tel. (boy see Tel.) Contact: JoeieTsang Email: bvcps_pyinfo@cn.bureauveritas.com

Hong Kong Bureau Veritas Hong Kong Limited L Decido Teade Centre, 1st Floor 2 Kai Hing Road, Kowloon Bay, Kowloon, Hong Kong Tet: 852-2331-0888 / Fax: 852-2331-0889 Sample Pick-up: 852-2331-0333 Toys, Premiums & Juvenile Products Contact: Eugene Wong Email: bvcps.toy@hk.bureauverites.com Inspections, Factory Audits & Social Audits Contact: Samuel Lau Email: bvcps.inspection@hk.bureauveritas.com

Bureau Veritas Hong Kong Limited L Unit 19, 13/F, Vanita Industrial Centre 21-33 Tei Lin Pei Road, Kwei Chung, N.T., HK Tel: 852-2418-1222 / Fax: 852-2485-1125 Softlines & Footwear Contact: Bonnic Chung Email: bvcps.softline@hk.bureauveritas.com Hardlines Contact: Priscilla Cheung Email: bucps.hardline@hk.bureauveritas.com Electrical & Electronics Contact: Christian Laudet Email: bycgs electrical@hk bureauveritas.com

India Bureau Ventas Consumer Products Services (I) Pvt. Ltd. L F-5, Sector 8, Noida Uttarpradesh, India 201 301 Tel: 00-91-120-2424-878 / Fex: 00-91-120-2424-880 Contact: Paromita Roy Email: paromita.roy@in.bureauveritas.com

Bureau Veritas Consumer Products Services (I) Pvt. Ltd. 1 1-2-3, Mereside Heights, Pai-Layout Old Madras Road, Benniganahalli Bangalore - 16, Karnatka, India Tel: 00-91-80-28531432 / Fax: 00-91-80-28531397 Contact: Paromita Roy Email: paromita.roy@in.bureauventas.com

Bureau Veritas Consumer Products Services (I) Pvt. Ltd. 4 212, Udyog Vihar, Phase – IV Gurgaon - 122002, Haryana, India Tel: 00-91-124-4548-100 / Fax: 00-91-124-2340-807 Contact: Paromita Roy Email: paromita.roy@in.bureauveritas.com

Bureau Veritas Consumer Products Services (I) Pvt. Ltd. 4 79/51, MRD Complex, Near Nesvalar Colony Bus Stop P.N. Road, Tirupur - 641602, Tamilnadu, India Tel: 0091-421-4308100 / Fax: 0091-421-4308105 Lab Contact: Sumantra Pal Email: sumantra.paul@in.bureauveritas.com Sales & Marketing: S.Palaniappan Email: s.palani@in.bureauveritas.com

Bureau Veritas Consumer Products Services (I) Pvt. Ltd. 9 50 Quiet Lands, Gashibowli, Hyderebad 500 032, Andhra Pradesh, India Tel: 0091-40-23000096 / Fax: 0091-40-23002964 Contact: S.Srinivasan Email: s.srinivasan@in.bureauveritas.com

Indonesia

PT. Bureau Veritas Consumer Products Services Indonesia L Jl. Cideng Timur No. 38, Gedung KKM, 2nd Floor Jakarta Pusat 10130, Indonesia Tel: 021-634-8877 / Fex: 021-634-8838 Contact: Darwi Tandiono Email: bvcps_jkt_general_inquiries@id.bureauveritas.com

Japan Synthetic Textile Inspection Institute Foundation ^L 4-4-20, Hongokutho, Nihonbashi, Chuo-Ku Tokyo 103-0021, Japan Tel: 81-3-3241-7319 / Fax: 81-3-3245-0773 Contact: Takuo Ishizuka Email: kikaku-info@kaken.or.jp

^{*}These locations are subject to changes. Please visit http://cps.bureauveritas.com for the most current listing of locations.



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Korea KOTITI (softines)

819-5 Yeoksam-1 Dong, Gangnam-Gu Seaul, Korea Tel: 822-3451-7000 / Fax: 822-3451-7172 Contact: Kyu Huh, Dr. Lee Myung-Hak Email: kyu_huh@kotti.re.kr, myunglee@kotti.re.kr

KEMTI L (hardines) 459-28 Gesen-Dong, Gumcheon-Gu Seoul, Korea Tel: 82-2102-2553 / Fex: 82-2-868-6903 Contact: Jeonghan Lee Email: komtri@chal.net

Pakistan

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Email: criselda.de-jesus@ph.bureauveritas.com or carmela.babor@ph.bureauveritas.com

Singapore
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Email: chanaki.de-costa@k.bureauveritas.com

Taiwan Bureau Veritas Consumer Products Services, Taiwan Branch L No. 37 Zhongyang S. Road, Sec. 2, Beitou, Taipei 112, Taiwan R.O.C. Tel: (886) 2 2895 3666 / Fex: (886) 2 2895 6999 Contact: Jane Lo Email: general.twncps@tw.bureauveritas.com

Advance Data Technology Corporation, L A Bureau Veritas Company No. 19, Hwa Ya 2nd Rd. Kwei Shan Hsiang Taoyuan Hsien 333, Taiwan (Hwa Ya Technology Park) Tel: +886-3-318-3232 / Fax : +886-3-211-5834 Contact: Mark Wang Email: service@adt.com.tw

Thailand

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Vietnam

Bureau Veritas Consumer Products Services Vietnam Limited 4 934 D3, D Street, Thanh My Loi Ward, Cat Lei Industrial Zone, District 2 Ho Chi Minh City, Vietnem Tel: 84-8-742-1605 / Fax: 84-8-742-1603 Contact: Mr. Dana Mason Email: dana.mason@hk.bureauveritas.com

KOTITI Vietnam Limited L (softines) Lot K, Road No. 2, Song Than Industrial Zone II, Di An District, Binh Duong Province, Vietnam Tel: 84-650 737 367 / Fax: 84-650 737 372 Contact: Dr. Young Kim or E.C. Park E-mail: young_kim@kotti.re.kr or ec_park@kotti.re.kr

Updated: 06/16/08

Bureau Veritas Consumer Products Services www.bureauveritas.com/cps

^{*}These locations are subject to changes. Please visit http://cps.bureauveritas.com for the most current listing of locations.





NOTE: MANDETORY FIELDS ARE MARKED WITH A DOUBLE ASTERIC (**) AND HIGHLIGHTED IN YELLOW. A HOLD MEMO WILL BE ISSUED TO THE SUBMITTER IF ANY MANDATORY INFORMATION IS MISSING AND TESTING WILL BE PLACED ON HOLD UNTIL EVERYTHING IS RECEIVED.

Please type or print in BLOCK			for each sample submitted)
	Toys and Hardline Prog	gram 🔲 Softlines	
Company Address	Dhana		Faulla
Contact Person	Phone		Fax No
Email address			
Special Instructions			
Report and invoice will state the ab			
	C	Contact Person	
		Contact Person	Fax No
Invoice To (Supplier): Company		Contact Person	Tel No.
		***************************************	Fax No.
Sample Description			
Style / Item #	Quantity	Country o	f Origin
**SKU # (List all)			**PO #
Product Phase Pre-tes	ting	——— ☐ Final F	Production
**Manufacturer/Factory	Δ ₁	gency (If Applicable)	
*If this is a Retest (first or second) and previous Test Report No(s). Corrective Action(s) Taken: 2nd Re-Test - Big Lots approve			
SERVICE REQUIRED:		SAMPLE RETURN	·
Regular Service		Self-Colle	
Next Two Days (40% surch	narge)		ect Fail Sample
Next Day (100% surcharge		Return Sa	amples *
Same Day (150% surcharg	,		ail Samples *
* Additional Charges apply (Pleas We apply for the above tests and agr forth in their current price list and up Date	ee that all testing will be carri	ed out subject to Burea	nu Veritas of charges and Turnaround Times as set in the back of this form.
	FOR LABOR	ATORY USE ONLY	
Date Rec'd	Rec'd By		em Due
Total Qty rec'd	Sample	No	
PLEASE SUBMIT YOUR SAMPLES T	O OUR APPROPRIATE LABOR	RATORY ADDRESS AS	BELOW:
Bureau Veritas - New York Office	Bureau Veritas Hong Kon	,	Bureau Veritas Hong Kong – Kwai Chung Office
(For Toys and Hardline items) 100 Northpointe Parkway, Buffalo, N 14228-1884 U.S.A. Tel: 716-505-3300 Fax: 716-505-3301	(For Toys, Child Care ite I.Y. 2/F Pacific Trade Cen Kowloon Bay, Hong Kong Tel: (852)23310222 Fax	tre, [°] 2 Kai Hing Road, _J .	(For Hardline, Softline, Electrical items) Unit 414, 4/F, Vanta Industrial Centre, 21-33 Tai Lin Pai Road, Kwai Chung, N.T., Hong Kong Tel: (852)24181222 Fax: (852)24806666

CONDITIONS OF TESTING

Bureau Veritas Consumer Products Services, Inc. ("BVCPS"), a Massachusetts corporation, or a subsidiary or affiliate of it, as identified in a written report (the "Test Report") issued by BVCPS or such subsidiary or affiliate (BVCPS or such subsidiary or affiliate being referred to herein as the "Company"), will conduct at the request of the Submitter ("Client"), the required tests specified on the reverse side of this Test Request Form in accordance with, and subject to, the following terms and conditions:

- 1. All orders for tests are subject to acceptance by the Company, and no order will constitute a binding commitment of the Company unless and until such order is accepted by it, as evidenced by the issuance of a written report ("Test Report") by the Company. The Test Report is issued solely by the Company, is intended for the exclusive use of Client and shall not be published, used for advertising purposes, copied or replicated for distribution to any other person or entity or otherwise publicly disclosed without the prior written consent of the Company. The Company shall not be liable for any loss or damage whatsoever resulting from the failure of the Company to provide its services within any time period for completion estimated by the Company. If Client anticipates using the Test Report in any legal proceeding, arbitration, dispute resolution forum or other proceeding, it shall so notify the Company prior to submitting the sample for testing.
- 2. The names, service marks, trademarks and copyrights of the Company and its affiliates, including the names "BUREAU VERITAS," "BUREAU VERITAS CONSUMER PRODUCTS SERVICES," "VERITAS" and "BVCPS", (collectively, the "Marks") are and shall remain the sole property of the Company or its affiliates and shall not be used by Client except solely to the extent that Client obtains the prior written approval of the Company and then only in the manner prescribed by the Company. Client shall not contest the validity of the Marks or take any action that might impair the value or goodwill associated with the Marks or the image or reputation of the Company or its affiliates.
- 3. The Test Report will set forth the findings of the Company solely with respect to the test samples identified therein and, unless specifically and expressly indicated in the Test Report, the results set forth in such Test Report will not be indicative or representative of the quality or characteristics of the lot from which a test sample will be taken. The Test Report will reflect the findings of the Company at the time of testing only, and the Company shall have no obligation to update the Test Report after its issuance. The Test Report will set forth the results of the tests performed by the Company based upon the written information provided to the Company as set forth in, or attached to, this Test Request Form. The Test Report will represent the entire understanding of the parties hereto with respect to the subject matter of the Test Report and no modification, variance, extrapolation or conclusion with respect thereto shall be permitted without the prior written consent of the Company.
- 4. The Company may, in its sole discretion, destroy samples which have been furnished to the Company for testing and which have not been destroyed in the course of testing. The Company may delegate the performance of all or a portion of the services contemplated hereunder to an affiliate, agent or subcontractor of the Company, and Client consents to such delegation.
- 5. Unless a shorter period is provided for on the reverse side of this Test Request Form, payment in full shall be due 30 days after the date of invoice. Client shall pay interest on any overdue amount from the due date until paid at an interest rate of 1.5% per month or, if less, the maximum rate permitted by law. The Company reserves the right, at any time and from time to time, to revoke any credit extended to Client. Client shall reimburse the Company for any costs it incurs in collecting past due amounts, including court costs and fees and expenses of attorneys and collection agencies. The Test Report may not be used or relied upon by Client if and for so long as Client fails to pay when due any invoice issued by the Company or any affiliate of it to Client or any affiliate or subsidiary of Client together with interest and penalties, if any, accrued thereon.
- 6. The Company may, from time to time, permit Client to access this Test Request Form, the Test Report and other communications by means of e-mail transmissions with the Company. Client acknowledges that any such transmission will not be encrypted and, hence, will not be confidential, that such transmissions may be read and intercepted by third parties and that the electronic version of a Test Request Form, Test Report or other communication could be modified inadvertently. The Company disclaims any and all responsibility or liability arising out of or in connection with e-mail transmissions of such information.
- 7. The Company represents and warrants solely to Client that the Test Report will be free of any material error or omission caused by the negligence of the Company. If Client desires to assert a claim for breach of the foregoing warranty, it must submit a claim to the Company within 60 days after the date of issuance of the Test Report to Client in a writing that sets forth with particularity the basis for such claim. If the Company determines that the claim is timely and that a breach of the foregoing warranty has occurred, then the Company, at its option, may either (a) re-perform the deficient test, without charge to Client, or (b) refund to Client, without interest, the fee paid to the Company for such Test Report. Client waives any and all claims for breach of the foregoing warranty, including, without limitation, claims that the Test Report is inaccurate, incomplete or misleading or that additional or different testing is required, unless and then only to the extent that Client submits a written claim to the Company within such 60-day period.
- 8. The Company shall, to the extent of the limitation of liability set forth in Section 9, indemnify Client against third party claims asserting a loss arising exclusively from the negligence of the Company, but only if Client (a) notifies the Company of the assertion of such claim immediately upon its being notified and (b) provides to the Company the option to assume the defense of such claim or participate in such defense.
- 9. EXCEPT TO THE EXTENT OF THE LIMITED WARRANTY SET FORTH IN SECTION 7 OR AS MAY OTHERWISE BE AGREED TO IN WRITING BY THE COMPANY AND NOTWITHSTANDING ANY PROVISION TO THE CONTRARY CONTAINED HEREIN OR IN ANY TEST REPORT, NO WARRANTY OR GUARANTEE, EXPRESS OR IMPLIED, INCLUDING ANY WARRANTY OF MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE OR USE, IS MADE. IN NO EVENT WHATSOEVER SHALL THE COMPANY BE LIABLE FOR ANY CONSEQUENTIAL, SPECIAL OR INCIDENTAL DAMAGES IN CONNECTION WITH OR ARISING OUT OF THE SERVICES PROVIDED BY THE COMPANY HEREUNDER, INCLUDING WITHOUT LIMITATION LOSS OF OR DAMAGE TO PROPERTY, LOSS OF INCOME, PROFIT OR USE, OR CLAIMS OR DEMANDS MADE AGAINST CLIENT OR ANY OTHER PERSON BY ANY THIRD PARTY IN CONNECTION WITH OR ARISING OUT OF THE SERVICES PROVIDED BY THE COMPANY HEREUNDER. NOTWITHSTANDING ANY PROVISION TO THE CONTRARY CONTAINED HEREIN, UNDER NO CIRCUMSTANCES WHATSOEVER SHALL THE LIBBILITY OF THE COMPANY IN RESPECT OF ANY CLAIM FOR LOSS, DAMAGE OR EXPENSE, OF WHATEVER NATURE OR MAGNITUDE, AND HOWEVER ARISING, EXCEED AN AMOUNT EQUAL TO THE AMOUNT OF THE FEES PAID TO THE COMPANY FOR THE SPECIFIC SERVICES WHICH GAVE RISE TO SUCH CLAIM.
- 10. The Company shall not be liable for any loss or damage resulting from any delay or failure in performance of its obligations hereunder resulting directly or indirectly from any cause that is beyond the control of the Company, including, but not limited to, fire, flood, tornado, hurricane or other acts of God, war, casualty, accident, embargo, governmental actions, orders of courts or tribunals, non-performance of third parties, strike, lock-out, or other difficulties with employees, inability timely to obtain labor, material, equipment or services through the Company's usual sources or delays of carriers. If any such event occurs, the Company may immediately cancel or suspend its performance hereunder without incurring any liability whatsoever to Client.
- 11. These Conditions of Testing shall be governed by, and construed in accordance with, the local laws of the country where the Company performs the tests or, in the case of tests performed in the United States of America, the laws of the Commonwealth of Massachusetts. Client waives the right to interpose any counterclaim or setoffs of any nature in any litigation arising hereunder. Any proceeding related to the subject matter hereof shall be brought, if at all, in the courts of the country where the Company performs the tests or, in the case of tests performed in the United States of America, in the courts of the Commonwealth of Massachusetts.
- 12. These Conditions of Testing are available at the website http://www.cps.bureauveritas.com. If there is a discrepancy between the terms and conditions set forth in these Conditions of Testing and the terms and conditions set forth (a) at such website; (b) in the Test Report, (c) on the reverse side of this Test Request Form or (d) in any other writing other than a master agreement which expressly (i) provides that its terms and conditions shall take precedence over these Conditions of Testing and (ii) cross-references this Section 12 or the equivalent section at such website, the Conditions of Testing set forth at such website shall govern and any inconsistent provision of this Test Request Form or other writing, as applicable, shall be disregarded. If no Conditions of Testing are set forth at such website, then the Conditions of Testing set forth herein shall govern.

B	ig Lots Fa	ctory Asses	sment Que	stionnaire	!				
Vendor Details: (This form is to be completed by the fa	ectond								
Vendor Name:									
Address:									
Telephone:			Fax:						
Name of Contact:			e-mail:						
Factory Details:									
Factory Name:									
Address:									
Telephone:			Fax:						
Name of Contact:			e-mail:						
Year Factory Established:			F		l				
Name of Factory Manager:	l		Fax:						
Telephone: Working Hours:			e-mail:						
Product:									
Total Workers:	Foreign wo		Local Workers:						
Total Male Workers:	Foreign workers: Total Female Workers:		LUCAI YVUINEIS.						
iotal male yyuineis.	iviai reillali	TYUINGIO.							
Domitory Address:									
(If Applicable)									
(Copposite of the Copposite of the Coppo									
CHILD LABOUR									
What is the age of the youngest worker in this facility?	Man minimum land madri		anding on the level laws						
Are there any restrictions for "young" workers? (between	. the minimum legal workin	ng age and 1% years old , dep	ending on the local law)						
If Yes , please give details.									
ii res , piedse give details.									
What procedures does this facility have in place to verify t	the age of the employees'	?							
with procedures does also identify flave in place to verify	are age or are employees	:							
What records does the facility obtain from job applicants b	efore hiring to determine :	age?							
Are age documents or records for all workers maintained	I at factory site?								
COMPENSATION									
JOHN LAGATION									
What is the legal minimum wage required at this factory?			Local Currency:						
What is the lowest wage paid at this factory for employees	6?								
		П Он	□ Electronic Paγ	Пон					
How are workers paid?	☐ Cash	□ Cheque		Other	(Please state	if other)			
How often are workers paid?	□ Weekly	□ Bi-weekly	☐ Monthly	□ Other	(Please state if other)				
How is the pay rate calculated?	□ Hour	☐ Piece Rate	□ Per Daγ	□ Other	(Please state if other)				
Are workers given work to take home?					□ Yes	□ No			
If Yes , how are wages for this work determined?]				
Is work time electronically documented by a timecard?					□ Yes	□ No			
If No , explain system by which hours worked are recor	ded?	'							
What records are used to calculate workers paid under a		licable)?			1				
Are there any deductions from employees' wages?	present to app				□ Yes	□ No			
What charges are deducted from employees' pay?									
Does this facility employ temporary workers?					□ Yes	□ No			

What allowances and benefits are provided to employees in this facility?					lity?					
Housing		Yes		l No		Sick leave	□ Ye	s 🗆 No		
Meals		Yes		I No		Emergency Leave	□ Ye	s 🗆 No		
Transportation		Yes		No		Maternity Leave	□ Ye	s □ No		
Health Care		Yes		No		Vacation	□ Ye	s 💹 No		
Child Care		Yes		No		Religious Leaves	□ Ye	s 🗆 No		
Other		Yes _		l No	If Yes , please give deta	ails				
			J							
Are any workers excluded	from receiv	ing benefits?							□ Yes _	L□ No
If Yes, please give details.										
Are benefits and /or allowances included in calculating the minimum wage? (E.v.	E.
If Yes, please give details.									⊥□ Yes _	⊥□ No _
Are there any incentive pla	and offered (ia honus)?							-	-
If Yes, please give details.	ano onerea q	i.c. borida):							⊥□ Yes _	⊥□ No _
Do you pay for overtime?							 □ N/A		□ Yes	
If Yes , how are overtime	wages calcu	ulated.					LINA			
										· · · · · · · · · · · · · · · · · · ·
DISCRIMINATION	1								,	
How are workers recruited	: :									
☐ Agent				Newspap	oer ad					
☐ Contract							1			
Do you restrict employment by age, ethnicity, religion, gender, sexual orientation or political affiliation?					if other , give details:	[
	it by age, eu	ir iicity,r eiigio	n, gen	uer,sexua	orientation or political alli	IIIduOrre				
If yes, please give details:										
DODUSTORY										
DORMITORY										
Do you provide residential	l housing for	workers?							□ Yes _	□ No _
If yes to the question above	e, please an	swer all que	stions I	below:						
							<u> </u>			
Number of buildings Average number of per					ople sleeping in a room					
No of employees residing i	in this dormit	ory								
Are sleeping quarters sepa	arated for ma	ale and fema	le?						□ Yes _	□ No
Are directions for evacuation plan posted in all sleeping quarters in employees native language?					de?					
					l language				□ Yes	□ No
Number of toilate for works	are:				1		Male	,.	Female:	
Number of toilets for workers: Number of shower's/ washing cubicles for employees:					Male		Female:			
					Widit		□ Yes	□ No		
Are kitchen facilities provided?							T 169 _	Tr 1/10		
Are there any curlews on employees?							□ Yes _	□ No		
If Yes, please give details.										
Do employees have unrestricted access to drinking water?								□ Yes	□ No	

EARLOS CARRELATE								
ENVIRONMENT	_							
Does the factory make any provision to protect the environment(e.g. wastewater management, air purification, hazardous material disposal)?	□ Yes	□ No						
Hazardoud Waste Disposal								
Does the factory have appropriate harzardous waste disposal procedure?	□ Yes	□ No						
Not Applicable.		 □ No						
Waste Water Treatment		140						
Is there waste water treatment facilities?	□ Yes	□ No						
Not Applicable.	□ Yes	No						
Air Emission								
	□ Yes	□ No						
Is there any equipment to control emission of air pollutant?								
Not Applicable.	□ Yes	No						
		L NO						
FACTORY COMPLIANCE WITH THE LOCAL LAW								
	le v	- ·						
Does the factory obtain current information on relevant local and national laws and regulation?	□ Yes	□ No						
FORCED LABOUR								
Are all workers voluntarily working in this facility?	□ Yes	□ No						
Are there any workers in this facility that are prisoners , have been assigned by the military , or any other branch of government?								
Are there any workers in this facility that are prisoners , have been assigned by the filminary , or any other branch of governments	□ Yes	□ No -						
Are workers free to leave the factory premises at any time?		L□ No _						
If No , please give details								
		1						
LIAD ACCURET AND ADVICE								
HARASSMENT AND ABUSE		1						
Are workers disciplined for misconduct or poor performance?		□ No						
What records are kept to document disciplinary measures?	□ Yes	□ No						
	······	· · · · · · · · · · · · · · · · · · ·						
Are there any procedures through which workers can appeal to disciplinary actions against them?	□ Yes	□ No						
HOURS OF LABOUR								
How many work shifts do you run in your facility?								
How many hours per work shift?								
(pls indicate start work timing, break timing and stop work timing):								
What is the maximum number of working hours per day?								
What is the maximum number of working hours per week?								
What is the average number of overtime hours worked per employee per month?								
Do workers have at least one day off in seven?	□ Yes	□ No _						
Do workers have time each day for a meal / breaks?	□ Yes	□ No _						
What are the break times? What is the duration of each break?								

HEALTH AND SAFETY									
Does the factory provide any of the below?									
Fire Safety Equipments / Systems							□ Yes	□ No	
First Aid / Health care?							□ Yes	_ No _	
Electrical & machinery safety?							□ Yes	□ No	
Completed by:									
Name:					Title:				
Signature:					Date:				