

Please read carefully before completing your documents inside.

1. COMMERCE CUSTOMS & TRANSPORATION ORDER FORM

Please complete the enclosed Customers Order Form, ensuring that all fields, including credit card information, are filled.

N.B. Goods cannot be returned to the U.S.A. after the show unless I.R.S./Tax, I.D./E.I.N. number is provided.

2. CANADA CUSTOMS DOCUMENTATION

(Photocopies are acceptable)

Enclosed is a Canada Customs invoice. <u>AN EXAMPLE FOR</u> COMPLETION PURPOSES IS ENCLOSED.

How to complete the Canada Customs Invoice:

- Fill in all information required on the Canada Customs Invoice, (Follow example enclosed.)
- B. Three (3) completed Canada Customs invoices must be attached to your Truck Bill of Lading, Air Waybill or Ocean Bill of Lading.
- C. The Bill of Lading or Air Waybill, all Customs documents and labels must be marked "NOTIFY COMMERCE FOR CUSTOMS CLEARANCE".
- D. Fax copies of the Canada Customs invoice, <u>ORDER FORM</u>, Truck Bill of Lading, Air Waybill, or Ocean Bill of Lading (and a copy of the FCC Information Sheet, if required).
- E. Retain a photocopy of all documents for your records and your on-site COMMERCE REPRESENTATIVE.

3A. ITEMS NOT OF U.S.A. ORIGIN AND TO BE RETURNED TO U.S.A. AFTER EVENT

VERY IMPORTANT:

For any items you are shipping to Canada that are <u>not of U.S.A.</u>, origin but have been shipped from the U.S.A., the U.S. Certificate of Registration (Form 4455) contained in this brochure is <u>required</u>. This form must accompany the shipment to Canada **and must be signed by U.S. Customs prior to coming to Canada**.

Complete the 4455 as per the example and MARK IN THE BODY OF YOUR BILL OF LADING OR AIR WAYBILL:

"CERTIFICATE OF REGISTRATION 4455 ATTACHED. GOODS MUST BE IDENTIFIED BY U.S. CUSTOMS AT PORT OF EXIT AND CERTIFIED COPIES TO BE GIVEN TO COMMERCE AT DESTINATION".

3B. ITEMS THAT EMIT OR ARE CAPABLE OF EMITTING RADIATION

Goods of this type: high voltage vacuum tube or switch, industrial X-ray systems, laser light show, microwave heating products, microwave ovens, CD players, etc., require an FDA Radiation From when they are returned to the U.S.A. Please complete the enclosed FDA Radiation Form to ensure your goods are not delayed upon return. Send the FDA Radiation Form with your Commerce Order Form.

3C. ITEMS THAT EMIT OR ARE CAPABLE OF EMITTING RADIO FREQUENCIES

Goods of this type: T.V.'s, communications equipment, computer equipment, monitors, microwaves, etc., require a Federal Communications Commission Statement when they are returned to the U.S.A. Please complete the enclosed F.C.C. Information Sheet to ensure your goods are not delayed upon return. Send the F.C.C. Form with your Commerce Order Form. Fax the F.C.C. Form to Commerce with a copy of the Order Form, the Canada Customs Invoice and your Bill of Lading or Air Waybill.

4. SHIPPING INSTRUCTIONS

Before shipping by Parcel Courier, i.e., UPS; call Commerce Customs Brokers for advice. We strongly suggest that all exhibitors DO NOT ship by parcel courier, or parcel post mail.

A) Exhibition goods shipped by common surface carrier should be scheduled to arrive and be available for Customs processing one week prior to show opening. Exceptions:

AIRFREIGHT — Schedule to arrive 3 days prior to show opening.

VAN LINE SHIPMENTS — May be sent direct to show site and should be scheduled to arrive on the appropriate move in day.

- B) All shipments **MUST BE SENT PREPAID.**
- C) For direct delivery shipments: all Truck Bills of Lading, Ocean Bills of Lading, Air Waybills and shipping labels should be addressed consigned to:

Exhibitor Name:

Booth:

Name of Event:

Location:

City/Prov: Zip Code:

Notify: COMMERCE for Customs Clearance.

5. PRIVATE/COMPANY VEHICLES

For goods being transported to Canada by: company, rented, personal or other non-Canada Customs Bonded method of transport; Canada Customs Invoices (and Certificate of Registration Form and F.C.C. Information Form if applicable), must accompany the shipment. You **MUST** notify us in advance with:

- A. Date and estimated time of crossing.
- B. Specific border crossing point.
- C. Total value and brief description of the goods.
- D. Driver's Name, Vehicle License Number and State shown on License Plate.
- E. A fax copy will be required for all appropriate forms including the Order Form

This information should be provided to our office one week prior to entering Canada.

6. AIRLINE HAND BAGGAGE

If you are bringing exhibit or commercial goods as airline baggage, in most cases "Formal" Customs clearance is required.

Contact our office at least one week in advance of your flight so arrangements can be made. Canada Customs Invoices (and Certificate of Registration Form and F.C.C. Information Form if applicable), must accompany the individual carrying the materials. A fac simile copy will be required for all appropriate forms including the Order Form.

7. SHOW SITE PROCEDURES

The show site has been declared a bonded area for the entire event. Under no circumstances are any goods to be removed without prior consent of COMMERCE. COMMERCE can provide the following services:

- Return of goods to your stated destination.
 Export Documents, Bill of Lading and labels supplied by COMMERCE.
- B) Arrange Customs clearance of those goods remaining in Canada. The applicable duties and taxes must be paid prior to removal from the show site.
- C) Arrange in-bond transfer of goods to be displayed at another Canada Customs recognized event.
- O) Coordinate under Customs supervision the destruction of non-returning items.
- Note: COMMERCE is not responsible for lost, stolen or damaged freight.
 All goods should be insured; prior to; during; and after the show.



T 905.673.5445
F 905.673.2574
Toll Free 1.888.827.7469

CUSTOMS AND TRANSPORTATION ORDER FORM

Complete and return all Custom and Shipping forms to the fax number indicated above. Forms must be received by The Commerce Trade Show Logistics Group Ltd. prior to the move in of your event.

Commerce can only process Customers, Transportation and/or related functions with a valid credit card. Individual Customs Order Forms must be used for multiple shipments. (photocopy as required)

		hibit will be shipped from: (city & int	ondod ompping dato)	
ABC COMPANY NEW YORK NEW YORK				
Name of Event	Fe	deral Tax I.D. #, Exporter Identification	on # or I.R.S. #	
XYZ ANNUAL EVENT	00	0-000000		
	oth # No	. of Pieces	Weight	
XYZ Convention Centre	149	2	2000	
Event Dates	Se	lected Carrier	•	
(M/D/YY)	X	YZ Transportation		
,	<u> </u>	,		
O Commence of the Commence of				
Our Company representative at the event will be: <u>John Doe</u>				
and can be reached at: ABC Hotel				
(name and telephone number	er of hotel - <i>for emergenc</i> y	y purposes only)		
At the completion of the quart all chipments will be chipped to				
At the completion of the event all shipments will be shipped to: Company Name			Contact Name	
ABC Company			John Doe	
Address	City		State/Province	
100 – 5 th Avenue Zip Code/Postal Code	Ne	ew York Fax #	New York	
Zip Code/Postal Code	1 el. #	Fax #		
00000	(212) 555-12	212 (21	2) 555-1212	
DILLING INCODMATION				
BILLING INFORMATION				
	rovided:			
Please indicate below method of payment you will be using for services p	rovided:			
	rovided: American Expres	ss 🗖		
Please indicate below method of payment you will be using for services p		ss 🗖	n Date: <i>01/10</i>	
Please indicate below method of payment you will be using for services possible. Master Card Visa 10045577046			n Date: <u>01/10</u> John Doe	
Please indicate below method of payment you will be using for services produced by Master Card Visa Credit Card Account Number: 12345678912		Expiratio		
Please indicate below method of payment you will be using for services proceed that the services		Expiratio Authorized Signature:	John Doe	
Please indicate below method of payment you will be using for services pure Master Card Visa Credit Card Account Number: 12345678912 Name of Cardholder: John Doe		Expiratio Authorized Signature:		
Please indicate below method of payment you will be using for services payment Card Wisa Credit Card Account Number: Name of Cardholder: Please forward copies of invoice to: Company name: ABC Company	American Expres	Expiratio Authorized Signature:	John Doe	
Please indicate below method of payment you will be using for services payment Card Wisa Credit Card Account Number: Name of Cardholder: Please forward copies of invoice to: Company name: ABC Company		Expiratio Authorized Signature:	John Doe	
Please indicate below method of payment you will be using for services payment Card Wisa Credit Card Account Number: Name of Cardholder: Please forward copies of invoice to: Company name: ABC Company	American Expres	Authorized Signature: Contact Name: Joh	John Doe	

Attention: The Commerce Trade Show Logistics Group Ltd. please accept this Order Form as your authority to effect <u>customs</u> <u>clearance of our shipment</u>. Commerce will process all Customs & related charges on the credit card that is listed above.

A statement will also be issued to the address listed above.

Any Carriers/Cartage companies that are providing a service for Commerce on behalf of the client listed above will receive a separate billing on the above credit card. This will only apply to Shows where the carriers/cartage company has a contracted agreement with Commerce.

Commerce will not be responsible for any goods delayed, lost, damaged, or stolen prior to, during or after the event.

Exhibitors are therefore urged to carry all -risk insurance covering their materials.



T 905.673.5445 F 905.673.2574

Toll Free 1.888.827.7469

CUSTOMS AND TRANSPORTATION ORDER FORM

Complete and return all Custom and Shipping forms to the fax number indicated above. Forms must be received by The Commerce Trade Show Logistics Group Ltd. prior to the move in of your event.

Commerce can only process Customers, Transportation and/or related functions with a valid credit card.

Individual Customs Order For	ms must be used for mu	Itiple ship		y as required)		
Exhibiting Company Name	Exhibit will be shipped from: (city & intended shipping date)					
Name of Event			Federal Tax I.D. #, Ex	porter Identificatior	n # or I.R.S. #	
Event Facility	Boo	oth #	No. of Pieces		Weight	
Event Dates			Selected Carrier			
Our Company representative at the event	t will be:					
and can be reached at:	<u> </u>					
and can be reached at:	(name and telephone number		r emergency purposes only)			
		01110101 101	emorganity purposes only)			
At the completion of the event all shipmer Company Name	its will be shipped to:					Contact Name
, , , , , , , , , , , , , , , , , , ,						
Address			City		State/Province	
Zip Code/Postal Code		Tel. #		Fax #		
p				1 227 11		
DILLING INCODMATION						
BILLING INFORMATION						
Please indicate below method of paymer	nt you will be using for services pr	rovided:				
Master Card \Box	Visa 🗖	Ameri	can Express 🔲			
Credit Card Account Number:				Expiration	Date:	
Name of Cardholder:			Authorized Si	ignature:		
Please forward copies of invoice to:						
Company name:			Contact Name) :		
-	_					
Address:	Cit	ty:			State/Province:	
-						
Zip Code/Postal Code:	Tel. #			Fax #		

Attention: The Commerce Trade Show Logistics Group Ltd. please accept this Order Form as your authority to effect <u>customs clearance of our shipment</u>. Commerce will process all Customs & related charges on the credit card that is listed above.

A statement will also be issued to the address listed above.

Any Carriers/Cartage companies that are providing a service for Commerce on behalf of the client listed above will receive a separate billing on the above credit card. This will only apply to Shows where the carriers/cartage company has a contracted agreement with Commerce.

Commerce will not be responsible for any goods delayed, lost, damaged, or stolen prior to, during or after the event.

Exhibitors are therefore urged to carry all -risk insurance covering their materials.



Page		
	Of	
	de	

							L		
Vendor (Name	and Address) / Vendeur (Nom	et adresse)					d'expédition dir	ecte vers le Canada	
Name and Address of Shipper				Date Shipped Must Be Shown 3. Other References (Include Purchaser's Order No.) Exporter's I.R.S./Tax I.D./E.I.N. Number Must Be Shown					
4. Consignee (Name and Address) / Destinataire (Nom et adresse)				5. Purchaser's Name and Address (if other than Consignee)					
Name of Exhibitor			١	Nom et adresse de	l'achet	eur (s'il diffère	du destinataire)	
City, Province Post	cility's Name, Street Address al Code		For	Display Purposes	Only				
Notify: COMMERC	E for Customs Clearance								
	ed company transaction? s compagnies sont liées entre e	lles?		Country of Tranship		•			
Yes	No			Country of Origin of Pays d'origine de	Goods	DIFFERE	ENT INCLUDES G NT ORIGINS ENT		
Oui \square	Non		n	narchandises			TITEMS IN 12. PÉDITION COMPRI	END DES	
				A., Japan Etc. t be Shown		MARCHA	NDISES D'ORIGINE	ES DIFFÉRENTES	
8. Transportation:	Give Mode and Place of Direct Sl	ninment to Canada		ondition of Sale and	d Terms		ISER LA PROVENA	ANCE EN 12.	
	siser mode et lieu d'expédition d		(i.	e. Sale, Consignme	ent Ship	ment, Leased			
Name of Carrier (Pi	lease Use "Canada Customs Bo	inded Carrier")		onditions de vente . ex. vente, expédi				archandises etc.)	
Name of Camer (if i	lease ose Canada Customs Do	inded Carrier)	No S	ale Involved		· ·		archandises, etc.)	
Place of Loading				urrency of Settlement of Be Shown	ent / De	vises du paie	ment		
11. No. of pkgs Nbre de colis	12. Specification of Commoditie	es (Kind of Packages,	Marks a	and Number,		uantity	Selling Pric	e / Prix de vente	
Note de coils	General Description and Cl Désignation des articles (I					State Unit) Juantité	14. Unit price	15. Total	
	description générale et ca				(Pré	ciser l'unité)	Price unitaire		
2 pcs	#1, #2 Wooden Crates Contai	ning Display Booth K	.D (Ba	ackwalls. Lights.		1	5,000.00	5,000.00	
•	Carpet, Graphics) and/or Des							•	
2 pcs	Cartons of Technical / Promot	tional Literature				5.000	.15	750.00	
1 pc	Cartons of Letter Openers					50	.50	25.00	
1 pc	Carton of Bottle Openers					50	1.00	50.00	
	Notify: COMMERCE for Custo	oms Clearance							
	of Pieces / Nombre total de Piece	es							
	er of Pieces Must Be Shown) to 17 are included on an attache	ed commercial invoice	check	this box	16	Total Waight	/ Poids total	17. Invoice Total	
	ements des zones 1 à 17 figuren					Total de la facture			
Commercial Invo	pice No, / N° de la facture comme	erciale			INCL		412 Kilos	Total Value Must Be Shown	
	e and Address (if other than Ver			20. Originator (I	Name a	ind Address)		Wast Be Griewii	
Nom et adresse	e de l'exportateur (s'il diffère de	vendeur)		Expéditeur Same as Si		ne (Nom et ad	dresse)		
Complete this Box	when the shipper is OTHER tha	n the Owner of these	Goods	Indicate Te	iippei lephoni	e Number and	d Contact Name		
(Contact Name, Co 21. Departmental Ru	mpany Name, Address, Telepho	one Number, Etc.)		(Person wh	o has k	nowledge of	this shipment)		
	érielle (s'il y a lieu)					• •			
				Si les zones	s 23 à 2	25 sont sans o	objet, cocher cet	te case 🔲	
	ld 17 indicate amount	24. If included in fie				25. Check (if			
	le total à la zone 17, préciser on charges, expenses and	Si compris dans	ie totai	à la zone 17, préci	ser		s'il y a lieu) Ity payments or s	ubsequent proceeds	
insurance fro Canada	m the place of direct shipment to			ges, expenses and	ont to		aid or payable by		
Les frais de	transport, dépenses et	Canada	iii tile pi	lace of direct shipme	ent to		u seront versés j	roduits ultérieurs ont par l'acheteur	
assurances à vers le Cana	partir du lieu d'expédition directe			rt, dépenses et lu lieu d'expédition d	liroeto				
	lua	vers le Cana		u neu a expedition a	llecte				
\$ (ii) Costs for con	nstruction, erection and assembly	\$				(ii) The r	ourchaser has su	pplied goods and	
incurred afte	er importation into Canada	(ii) Amounts for		sions other than bu	ıying	servi	ces for use in the	production of these	
	e construction, de montage et es après importation au Canada	commissions Les commiss		tres que celles vers	sées	good L'ach	s ieteur a fourni de	s biens ou des	
_	, , , , , , , , , , , , , , , , , , , ,	pour l'achar		,		servi	ces pour la produ		
\$(iii) Export packi	 ng	\$(iii) Export packi	ng			merc	handises.		
	emballage d/exportation			age d/exportation					
\$\$									



Page		
	Of	
	de	

1.	Vendor (Name and Address) / Vendeur (Nom et ad	dresse)	2. [Date of Di	rect Shipm	ent to	Canada /Date	d'expédition din	ecte vers le	Canada
			3. (Other Ref	erences (Ir	nclude	Purchaser's	Order No.)		
4.	Consignee (Name and Address) / Destinataire (No	nm et adresse)						than Consignee du destinataire		
VII.	1 is this a related company transaction?		6. C	Country o	f Transhipr	ment /	Pays de trans	sbordement		
	Est-ce que les compagnies sont liées entre elles?	′								
	Yes No Non Non		F	Country o Pays d'ori marchand		Goods	DIFFERE AGAINST S'IL L'EXF MARCHA	ENT INCLUDES G NT ORIGINS ENT ITEMS IN 12. PÉDITION COMPRE NDISES D'ORIGINE ISER LA PROVENA	ER ORIGINS END DES ES DIFFÉREN	
8.	Transportation: Give Mode and Place of Direct Shipm Transport: Préciser mode et lieu d'expédition direc		(i. Ce (p	e. Sale, Conditions o. ex. ven	Consignme de vente e te, expédit	nt Ship et mod tion er	J	ement n, location de ma	archandises	s, etc.)
				-		nt / De	evises du paie			
11.	No. of pkgs Nbre de colis No. of pkgs Nbre de colis General Description and Chara				ber,		Quantity State Unit)	Selling Price		
	Désignation des articles (Natu	ature des colis, marques et numéros, Quanti				Juantité ciser l'unité)	14. Unit price Price unitaire	15. T	otal	
XI.	Total number of Pieces / Nombre total de Pieces									
	lf any of fields 1 to 17 are included on an attached on Si les renseignements des zones 1 à 17 figurent sur					6. Total Weight / Poids total		17. Invoic		
	Commercial Invoice No, / N° de la facture commercia		/ G.G [Net		Gross /		
19.	Exporter's Name and Address (if other than Vendor) Nom et adresse de l'exportateur (s'il diffère de vend)		_			and Address) ne (Nom et ad	dresse)		
	Departmental Ruling (if applicable) Décision ministérielle (s'il y a lieu)			22. If 1	fields 23 to	25 ar	e not applicab	le, check this bo	×	7
	Decision immoterione (e ii y a nea)			Si	les zones	23 à 2	25 sont sans o	bjet, cocher cett	e case	
	Si compris dans le total à la zone 17, préciser (i) Transportation charges, expenses and insurance from the place of direct shipment to Canada Les frais de transport, dépenses et assurances à partir du lieu d'expédition directe vers le Canada \$	Si compris dans (i) Transportatio insurance from Canada Les frais de t	eld 17 indicate amount le total à la zone 17, préciser on charges, expenses and om the place of direct shipment to transport, dépenses et à partir du lieu d'expédition directe ada			nt to	(i) Royalty payments or subseque are paid or payable by the pur Des redevances ou produits ui été ou seront versés par l'ache		y the purcha roduits ultér	aser rieurs ont
	(ii) Costs for construction, erection and assembly incurred after importation into Canada Les coûts de construction, de montage et d'assemblages après importation au Canada \$ (iii) Export packing	(ii) Amounts for a commissions Les commiss pour l'achar (iii) Export packir	ions au		•		servio good L'ach servio	ourchaser has su ces for use in the s leteur a fourni de ces pour la produ handises.	production s biens ou o	of these
	Le coût de l'emballage d/exportation	Le coût de l'é		age d/exp	ortation					
	\$	\$								



Certificate of Registration Form 4455

	DO NOT WRITE IN	SHADED ARE	EAS			
MAKE COPIES AND	ATTACH THREE (3)	COPIES TO Y	OUR BILL OF	LADING	Form Approv	red OMB No. 48 -R0247
			COMPLETE ARE NOT OF	_		
VIA (Carrier)	mnany Airlina ata		B/L or INSURED		DATE	Must be Shown
NAME, ADDRESS AND ZIP (FORM IS TO BE M		if Available	ARTICLES EXP	
(If Applicable)	OODE TO WINOTI CERTIFIED	TOTAL TO DE IVI	AILLE		ANTIGLES EXP	UNIED FUN.
ABC Company				ALTERATIO	N *	□ PROCESSING *
3405 American Drive, Unit Mississauga, Ontario L4V				REPAIR * USE ABRO	nan	☐ OTHER, (specify)
•				REPLACE		- EXHIBITION
						ne cost or value of alterations, repairs
		LICT	ARTICLES EXPO)TEN	or process	ing abroad is subject to Customs duty.
NUMBER PACKAGES	KIND OF PACKAGES	DESCRIPTION	MULIOLES EXPUI	וובט		
NOWIDERT FACINALES	KIND OF FACINALES					
3 Crates (1 -3)	Wooden Crates as Addressed	Crate #2 - (Make	e / Brand Name) M	odel A - Com	mputer Serial #123 - Ma nputer Serial #456 - Ma nputer Serial #789 - Ma TO: (Cons	nde in Hong Kong nde in Taiwan
		ABC Company 123 Main Street New York, New	York 10001			oany ery Show, Booth #333 ddress of Event Facility
TYPE NAME	OWNER OR AGENT (Print or		_ TEL:()		DATE:
		THE ABOVE	-DESCRIBED ARTI	CLES WERE	:	
	EXAMINED				LADEN under m	y supervision
DATE	PORT		DATE			PORT
SIGNATURE OF CUSTOMS	OFFICER		SIGNA	ATURE OF C	USTOMS OFFICER	
		CER	TIFICATE ON RET	JRN		
Duty-free entry is claimed (see reverse if needed)	I for the described articles	as having been ex	xported without be	enefit of dra	wback and are returne	d unchanged except as noted:
	eeds the available space per Attached". Please at		• • • • • • • • • • • • • • • • • • • •			on this form
	mark the following in the S. Customs prior to exp				-	ion Form 4455 Attached . Goods at show site.
SIGNATURE OF IMPORTER (PI	rint or Type <u>and</u> Sign)				DATE	
	NOTE: Certifying offi	cers shall draw lir	nes through all un	used spaces	s with ink or indelible p	encil.



Certificate of Registration Form 4455

	DO NOT WR	RITE IN SHADED AR	EAS						
MAKE COPIES AND	ATTACH THRE	EE (3) COPIES TO Y	OUR BILL	OF L	_ADING			ed OM	B No. 48 -R0247
					WHEN GOOD U.S.A. ORIG).		
VIA <i>(Carrier)</i>			B/L or INSU	RED N	10.	DA	ΙΤΕ		
NAME, ADDRESS AND ZIP ((If Applicable)	CODE TO WHICH CE	RTIFIED FORM IS TO BE M	I MAILED			ARTI	CLES EXP	ORTED) FOR:
(п Аррисаліс)					ALTERATION * REPAIR * USE ABROAD REPLACEMENT			ne cost	□ PROCESSING * □ OTHER, (specify) EXHIBITION or value of alterations, repairs oad is subject to Customs duty.
NUMBER PACKAGES	KIND OF PACKA		T ARTICLES EX	(POR	TED				
TION SERVICES		EXPORTER:					ТО: (Cons	ignee)	DATE:
TYPE NAME		Print or Type <u>and</u> Sign)	- Te	EL: ()				
		THE ABOVE	-DESCRIBED A	ARTIC	CLES WERE:				
	EXAMINED					LADE	N under m	y super	rvision
DATE	POF	RT	D.	ATE				PORT	
SIGNATURE OF CUSTOMS	OFFICER		S	IGNA	TURE OF CUSTO	MS OFF	ICER		
			RTIFICATE ON I						
Duty-free entry is claimed (see reverse if needed)			exported withou	ut be			re returne	d uncl	hanged except as noted:
SIGNATURE OF IMPORTER (Pr		n) ying officers shall draw li	ines through al	ll unu	DATE		ndelible r	encil.	





FEDERAL COMMUNICATIONS COMMISSION Washington, DC 20554

Approved by OMB 3060-0059

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

PART I - All Blocks MUST Be Comp	pleted						
Date of Entry	Entry Number	Port of Entry	Harmonized Tariff Number			Quantity of item (not container)	
						1	
	U	o not complete shaded a	rea				
Model/Type Name or Model/type N	Number	Trade Name	** FCC ID Numb	per	Descrip	tion of Equipment	
LaserJet 4		HEWLETT PACKARD		B94C2001A		PRINTER	
Manufacturer's Name and Address	<u> </u>		Importer's Name	e and Address			
			ABC CO				
HEW	LETT PACKA	RD	123 MAIN STREET				
1131	1 CHINDEN B	I VD				T = .	
			Signature of Impo	orter or Consignee		Date	
			D	OBERT SMITH		JAN	
Consigneeís Name and Address			"	ODENT SWITT		JAN	
9			Print or Type Mar	me of Importer or Consigned			
SAM	IE AS IMPORT	ΓER	Print or Type Name of Importer or Consignee				
			DODEDT CM	ITI I			
	ROBERT SMITH						
WA BANK	10. 4	 					
WARNIN	lG: Any perso	on who knowingly makes	ataise declai	ration may be fined	i not m	ore tnan	

\$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.)

X	1. An equipment authorization has been issued by the FCC.
	2. An equipment authorization is not required but the equipment complies with FCC technical requirements.
	The described equipment is being imported in limited quantities for testign 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.
	6. The described equipment is being imported for use exclusively by the U.S. Government.
	Three or fewer radio receivers, computers, or other unintentional radiators as defined in Part 15 of the FCC Rules, are being imported for an individuals personal use and are not irtended for sale.
	8. The described equipment is being imported for repair and will not be offered for sale or otherwise marketed.

- * If shipping multiple pieces of equipment with the same Model # and FCC ID #, complete only one form.
- ** If no FCC ID # appears on equipment, leave blank.

FCC Form 740



FEDERAL COMMUNICATIONS COMMISSION Washington, DC 20554

Approved by OMB 3060-0059

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

PART I – All Blocks MUST Be Com	pleted						
Date of Entry	Entry Number		Port of Entry		Harmonized Tariff Number		Quantity of item (not container)
		o not comple	te shaded a	rea			
Model/Type Name or Model/type N	Number	Trade Name		** FCC ID Numb	per	Descript	tion of Equipment
						l	
						<u> </u>	
Manufacturer's Name and Address	3			Importer's Name	and Address		
				Ciamatana at lasa			l Dete
				Signature of impo	orter or Consignee		Date
Consigneeís Name and Address							
				Print or Type Nar	me of Importer or Consignee		
WARNIN	G: Any perso	n who knowi	ngly makes	afalse declai	ration may be fined	not m	ore than
			• •		th, pursuant to 18 U		
ΨZ	JU,UUU UI IIII	prisoneu not	illole alali 3	ycars, or nor	ııı, pursuallı tu 10 u	.3.0. 1	UU I .

PART II — With Regard to the Importation of the Described Radio Frequency Device(s), I DECLARE THAT: (Place an "X" in only one box.)

An equipment authorization has been issued by the FCC.
2. An equipment authorization is not required but the equipment complies with FCC technical requirements.
 The described equipment is being imported in limited quantities for testin and evaluation for compliance with technical requirements or marketing suitability. The equipment will not be offered for sale or otherwise marketed. (See instructions.)
 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.
6. The described equipment is being imported for use exclusively by the U.S. Government.
Three or fewer radio receivers, computers, or other unintentional radiators as defined in Part 15 of the FCC Rules, are being imported for an individuals 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.

- * If shipping multiple pieces of equipment with the same Model # and FCC ID #, complete only one form.
- ** If no FCC ID # appears on equipment, leave blank.



Form FD 2877

DEPARTMENT OF HEALTH, EDUCATION, AND WELFARE PUBLIC HEALTH SERVICE FOOD AND DRUG ADMINISTRATION

DECLADATION FOR PRODUCTS SUBJECT TO

Form Approved OMB No. 57-R0120

INSTRUCTIONS

- 1. Type or print with ball point pen. (One form per model #)
- 2. Complete one copy for U.S. Customs Commerce

PORT OF ENTRY For Customs Use Only For Customs Use Only For Customs Use Only PRODUCT IDENTIFICATION NAME AND ADDRESS OF MANUFACTURER: MATSUI JAPAN NAME AND ADDRESS OF IMPORTER OF RECORD: ABC COMPANY 123 Main Street New York, New York 10001 ULTIMATE CONSIGNEE (If not Importer of Record) OUANTITY TYPE COLOUR MONITOR BRAND NAME PANASONIC E1-16611 FOR X-RAY, LIST APPROPRIATE SYSTEMS OR COMPONENT CATEGORY COMPLETE ONLY WHEN SHIPPING X -RAY EQUIPMENT A That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: A That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is afficiate diverse or training. An exception for these products has been or will be requested of the Director of the PAB Juneau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the PAB Juneau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standard has been or will be filed with the Food and Drug Administration, that the products are in comp	RADIATION CONTROL STANDARDS	3. Attach all completed	3. Attach all completed copies to the Canada Customs Invoice					
PRODUCT DENTIFICATION NAME AND ADDRESS OF MANUFACTURER: MATSUI JAPAN NAME AND ADDRESS OF IMPORTER OF RECORD: ABC COMPANY 123 Main Street New York, New York 10001 ULTIMATE CONSGNEE (If not Importer of Record) OUANTITY TYPE COLOUR MONITOR PANASONIC COMPLETE ONLY WHEN SHIPPING X - RAY ECUIPMENT AFFIRMATION (Check appropriate statement and sign) IWE hereby declare: A That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: A That the electronic products identified above ocomply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. D. That the electronic products identified above ocomply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 3696 () (42 U.S. C. 326) of the Radiation Control for Health and Safety Act well will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filted with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce unit in indiffication is received from the Food and Drug Administration, that the products are in compliance with applicable standa	DISTRICT / PORT DIRECTOR OF CUSTOMS							
NAME AND ADDRESS OF MANUFACTURER: MATSUI JAPAN NAME AND ADDRESS OF IMPORTER OF RECORD: ABC COMPANY 123 Main Street New York, New York 10001 ULTIMATE CONSIGNEE (if not Importer of Record) OUANTITY TYPE COLOUR MONITOR BRAND NAME PANASONIC EI-1611 FOR X-RAY, LIST APPROPRIATE SYSTEMS OR COMPONENT CATEGORY COMPLETE ONLY WHEN SHIPPING X ATA THAT the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product to this effect is affixed to each product. C. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training, An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 3608 (b) (42 U.S. C. 263) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with applicable standards.	PORT OF ENTRY	ENTRY NO.	DATE					
NAME AND ADDRESS OF MANUFACTURER: MATSUI JAPAN NAME AND ADDRESS OF IMPORTER OF RECORD: ABC COMPANY 123 Main Street New York, New York 10001 ULTIMATE CONSIGNEE (If not Importer of Record) QUANTITY 1 TYPE COLOUR MONITOR BRAND NAME PANASONIC EI-1611 FOR X - RAY, LIST APPROPRIATE SYSTEMS OR COMPONENT CATEGORY COMPLETE ONLY WHEN SHIPPING X -RAY EQUIPMENT AFFIRMATION (Check appropriate statement and sign) LIWE hereby declare: A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: A That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Durau of Radiological Health in accord with Section 3680 (b) (42 U.S.C. 263) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the pr	For Customs Use Only	For Customs Use Only	For Customs Use Only					
NAME AND ADDRESS OF IMPORTER OF RECORD: ABC COMPANY 123 Main Street New York, New York 10001 ULTIMATE CONSGINEE (If not importer of Record) OUANTITY 1 TYPE OLOUR MONITOR BRAND NAME PANASONIC EI-1611 FOR X-RAY, LIST APPROPRIATE SYSTEMS OR COMPONENT CATEGORY COMPLETE ONLY WHEN SHIPPING X -RAY EQUIPMENT AFFIRMATION (Check appropriate statement and sign) IWE hereby declare: A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training, An exception for these products has been or will be requested of the Director of the FDA Eurose of Radiological Health in accord with Section 3680 (b) (42 US.C. 263) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 11010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filled with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicab								
NAME AND ADDRESS OF IMPORTER OF RECORD: ABC COMPANY 123 Main Street New York, New York 10001 ULTIMATE CONSIGNEE (If not Importer of Record) ULTIMATE CONSIGNEE (If not Importer of Record) ULTIMATE CONSIGNEE (If not Importer of Record) OUANITIY TYPE COLOUR MONITOR BRAND NAME PANASONIC EI-1611 FOR X - RAY, LIST APPROPRIATE SYSTEMS OR COMPONENT CATEGORY COMPLETE ONLY WHEN SHIPPING X - RAY EQUIPMENT AFFIRMATION (Check appropriate statement and sign) IWE hereby declare: A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: X B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 3608 (b) (42 U.S.C. 263) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into commerce with applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is								
ABC COMPANY 123 Main Street New York, New York 10001 ULTIMATE CONSIGNEE (If not Importer of Record) QUANTITY 1 TYPE COLOUR MONITOR BRAND NAME PANASONIC EI-1611 FOR X - RAY, LIST APPROPRIATE SYSTEMS OR COMPONENT CATEGORY COMPLETE ONLY WHEN SHIPPING X - RAY EQUIPMENT AFFIRMATION (Check appropriate statement and sign) I/WE hereby declare: A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: A. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filled with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.								
ULTIMATE CONSIGNEE (If not Importer of Record) QUANTITY TYPE COLOUR MONITOR BRAND NAME PANASONIC EI-1611 FOR X - RAY, LIST APPROPRIATE SYSTEMS OR COMPONENT CATEGORY COMPLETE ONLY WHEN SHIPPING X - RAY EQUIPMENT AFFIRMATION (Check appropriate statement and sign) I/WE hereby declare: A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 3608 (b) (42 U.S.C. 263) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the Administration in accordance with 21 CFR 1021. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.	NAME AND ADDRESS OF IMPORTER OF RECORD:							
ULTIMATE CONSGNEE (If not Importer of Record) QUANTITY TYPE COLOUR MONITOR PANASONIC 1 COLOUR MONITOR FOR X - RAY, LIST APPROPRIATE SYSTEMS OR COMPONENT CATEGORY COMPLETE ONLY WHEN SHIPPING X - RAY EQUIPMENT AFFIRMATION (Check appropriate statement and sign) IWE hereby declare: A That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: X B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263)) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to briging the product into compliance with the applicable standard has been or will be filled with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.								
QUANTITY TYPE COLOUR MONITOR BRAND NAME PANASONIC EI-1611 FOR X - RAY, LIST APPROPRIATE SYSTEMS OR COMPONENT CATEGORY COMPLETE ONLY WHEN SHIPPING X -RAY EQUIPMENT AFFIRMATION (Check appropriate statement and sign) I/WE hereby declare: A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training, An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to brigh the production tho compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.	New							
TOUR MONITOR PANASONIC EI-1611 FOR X - RAY, LIST APPROPRIATE SYSTEMS OR COMPONENT CATEGORY COMPLETE ONLY WHEN SHIPPING X -RAY EQUIPMENT AFFIRMATION (Check appropriate statement and sign) IWE hereby declare: A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263j) of the Radiotion Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.		Tork, now Tork Todo T						
TOUR MONITOR PANASONIC EI-1611 FOR X - RAY, LIST APPROPRIATE SYSTEMS OR COMPONENT CATEGORY COMPLETE ONLY WHEN SHIPPING X -RAY EQUIPMENT AFFIRMATION (Check appropriate statement and sign) IWE hereby declare: A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263j) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.			Lugher vo					
AFFIRMATION (Check appropriate statement and sign) I/WE hereby declare: A That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263)) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filled with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standard.								
AFFIRMATION (Check appropriate statement and sign) I/WE hereby declare: A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263j) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.								
AFFIRMATION (Check appropriate statement and sign) I/WE hereby declare: A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263j) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.								
A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 3608 (b) (42 U.S.C. 263j) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards in accordance with applicable standards.	COMPLETE ONLY WHEN SHIPPING X -RAY EQUIPMENT							
A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 3608 (b) (42 U.S.C. 263j) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards in accordance with applicable standards.								
A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard. Date of Manufacture: B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263j) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.	AFFIRMATION	(Check appropriate statement and sign)						
Date of Manufacture: B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263j) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.	I/WE hereby declare:							
B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263j) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.								
Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or lab el to this effect is affixed to each product. C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263j) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.	Date of Manufacture:							
Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263j) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision. D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.	Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label							
Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.	Administration Rules 21 CFR 1010 but are be training. An exception for these products has accord with Section 360B (b) (42 U.S.C. 263 commerce, and when their mission is complete.)	eing imported for the purpose of rese s been or will be requested of the Dir j) of the Radiation Control for Health	earch, investigations, studies, demonstrations or rector of the FDA Bureau of Radiological Health in and Safety Act. They will not be introduced into					
Signature of Importer of Record Robert Smith	Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug							
	Signature of Importer of Record Robert Smith							



Form FD 2877

DEPARTMENT OF HEALTH, EDUCATION, AND WELFARE PUBLIC HEALTH SERVICE FOOD AND DRUG ADMINISTRATION

DECLARATION FOR PRODUCTS SUBJECT TO

Form Approved OMB No. 57-R0120

INSTRUCTIONS

- 1. Type or print with ball point pen. (One form per model #)
- 2. Complete one copy for U.S. Customs Commerce

RADIATIO	N CONTROL STANDARDS		3. Attach all completed copies to the Canada Customs invoice			
DISTRICT / PORT DIRECTOR OF CUSTOMS						
PORT OF ENTRY		ENTRY N	0.	DATE		
PRODUCT IDENTIFICATION						
NAME AND ADDRESS OF MAN						
NAME AND ADDRESS OF IMPORE ULTIMATE CONSIGNEE (If not						
OLITIVIATE CONSIGNEE (II TIOL	Importer of necordy					
QUANTITY	TYPE		BRAND NAME		MODEL NO.	
FOR X - RAY, LIST APPROPRIATE SYSTEMS OR COMPONENT CATEGORY						
101/5		TION (Check appl	ropriate statement and sign)			
I/WE hereby declare: A. That the electronic products identified above were manufactured prior to the date of any applicable electronic product performance standard.						
	Date of Manufacture:					
B. That the electronic products identified above comply with the performance Standards prescribed in Food and Drug Administration Rules 21 CFR 1010 which are applicable at date of manufacture and that a certificate in the form of a tag or label to this effect is affixed to each product.						
C. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but are being imported for the purpose of research, investigations, studies, demonstrations or training. An exception for these products has been or will be requested of the Director of the FDA Bureau of Radiological Health in accord with Section 360B (b) (42 U.S.C. 263j) of the Radiation Control for Health and Safety Act. They will not be introduced into commerce, and when their mission is completed they will be destroyed or exported under United States Customs Service supervision.						
D. That the electronic products identified above do no comply with the performance standards prescribed in Food and Drug Administration Rules 21 CFR 1010 but that a timely and adequate petition for permission to bring the product into compliance with the applicable standard has been or will be filed with the Food and Drug Administration in accordance with 21 CFR 100.21. These products will remain under bond and not be introduced into commerce until notification is received from the Food and Drug Administration, that the products are in compliance with applicable standards.						
Signature of	Importer of Record					