



Medicines & Healthcare products  
Regulatory Agency

# Certificates of Free Sale

## Reference Guide

Please do not print this document. View online only to ensure you have the latest version.

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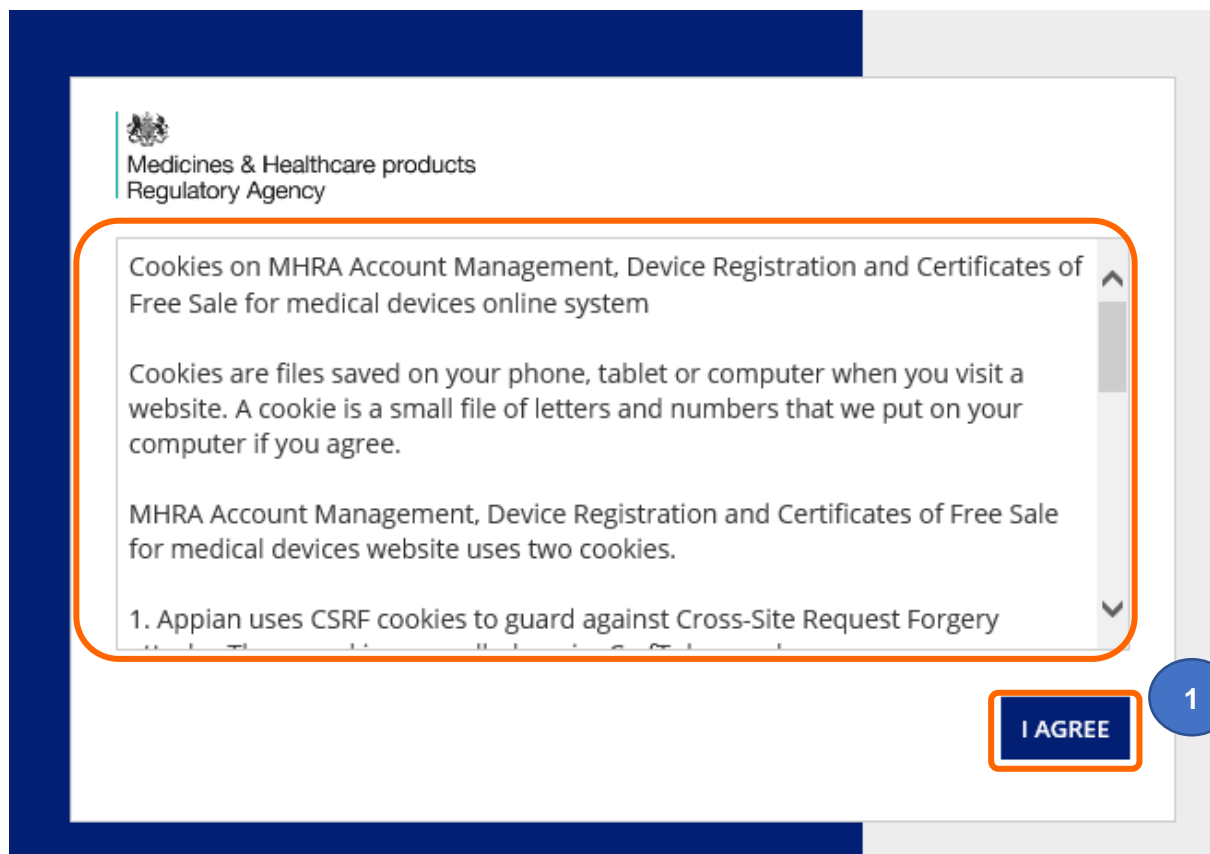
## Logging in

Access MHRA Agency Services for Device Registrations and Certificates of Free Sale for medical devices.

### Agree to Cookie Policy

Before accessing MHRA Agency Services, you will need to agree to our Cookie Policy. Please read the Cookie Policy and only use MHRA Agency services if you agree.

1. When you have read the Cookie Policy, **click** the 'I Agree' button.



## Username and Password

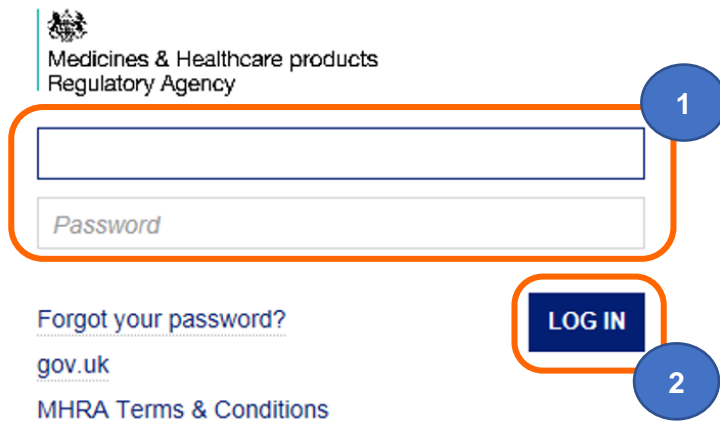
Once your Account request has been accepted by MHRA, two emails will be sent to the email address you entered in your account request application:


1. A welcome email with subject line **Account creation – outcome**, from email address [no-reply@mhra.gov.uk](mailto:no-reply@mhra.gov.uk) with instructions on initial actions to take in the registration system
2. A separate email with subject line **MHRA Portal account creation** from email address [admin@mhrabpm.appiancloud.com](mailto:admin@mhrabpm.appiancloud.com) containing your username (usually firstname.lastname), a temporary password and a link to the system

Please log in for the first time on a laptop or PC not a mobile or tablet. If you have not received the emails, please check your Junk/Spam folder. You will be asked to change the [password](#) to one of your choosing.

If the welcome email or the username and temporary password email have not been received this is usually due to your system blocking the originating email address. Please add the above email addresses to your **safe senders** list, usually via settings in your email system and email [device.registrations@mhra.gov.uk](mailto:device.registrations@mhra.gov.uk) to obtain your username and further instructions.

1. On the [log in](#) page, **enter** the [details](#) sent to you by email (it is preferable for you to [copy and paste](#) your details [into the boxes provided](#)).
2. **Click** the 'Log in' button.



  
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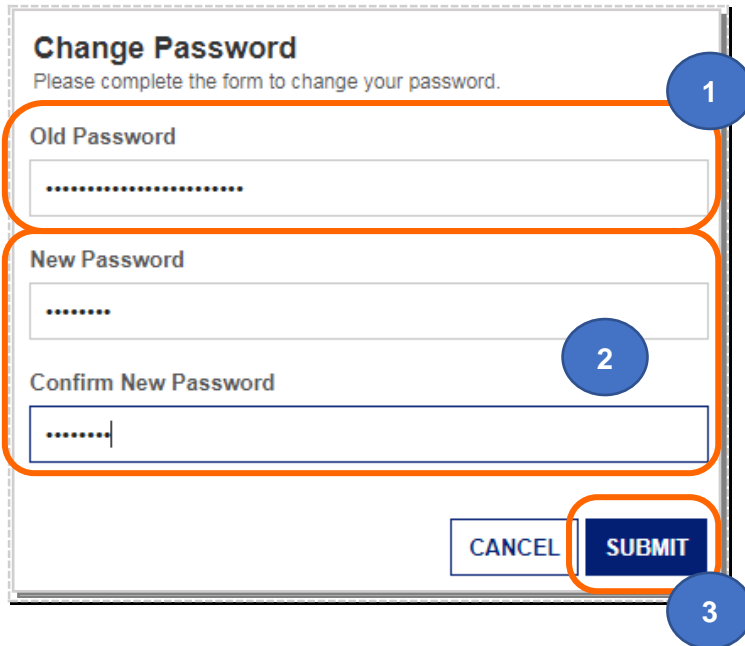
[Forgot your password?](#)

[gov.uk](#)

[MHRA Terms & Conditions](#)

**LOG IN**

## New Users > Change temporary password



**Change Password**  
Please complete the form to change your password.

1. Old Password

2. New Password

Confirm New Password

CANCEL SUBMIT

3.

1. **Copy** and **paste** the temporary password (long password with multiple characters) sent to you via email into the old password box.
2. **Enter** a **password** of your choice into the new password and confirmation **boxes**.
3. **Click** on **Submit**.  
You will be able to use the password you entered from now on.

## Forgot password > resets

1. On the [log in](#) page, **click** the 'Forgot your password' link.
2. **Enter** your **username** (usually firstname.lastname – not your email address).
3. **Click** the 'Send email' button.

You will be sent an email containing a link. Please check your Junk/Spam folder. **Click** on the link and follow the instructions to change your password. Please do this on a Laptop/PC not on a mobile/tablet.

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1

**Forgot your password?**

**LOG IN**

[gov.uk](#)

[MHRA Terms & Conditions](#)





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## Forgot Password

2

Username

Enter your username and click 'Send Email'. An email will be sent to the email address associated with your user account. Follow the link in the email to reset your password.

[Back to sign-in page](#)

3

SEND EMAIL

## Certificates of Free Sale (CFS) Service

This service enables UK based manufacturers, UK Responsible Persons and Authorised Representatives in Northern Ireland to apply for Certificates of Free Sale for medical device products that have been registered with MHRA.

Please follow all the instructions in the [Device Registration Reference Guide](#) to register devices for your organisation or manufacturers you represent.

Please note the important information in the above guide on adding product (model/version Catalogue/Reference and UDI DI etc.) data – only the [Medical Device Name, Model/Version](#) and [Catalogue/Reference, Basic UDI DI and UKCA or CE Certificate Reference No.](#) data that you enter will appear on the Certificate of Free Sale.

*in vitro diagnostic* medical devices for performance evaluation cannot be included on CFS orders.

It is your responsibility to check with the receiving country that they will accept the CFS **before** you make payment for your order. You will be given the option to download a preview sample that you can email to the country. Please note the formatting on the preview sample may not match the final certificate formatting.

CFS order applications are **non-refundable** – please see our terms and conditions. You must preview the CFS Certificate and Schedule **before** making payment to ensure it meets your needs and **before** you pay.

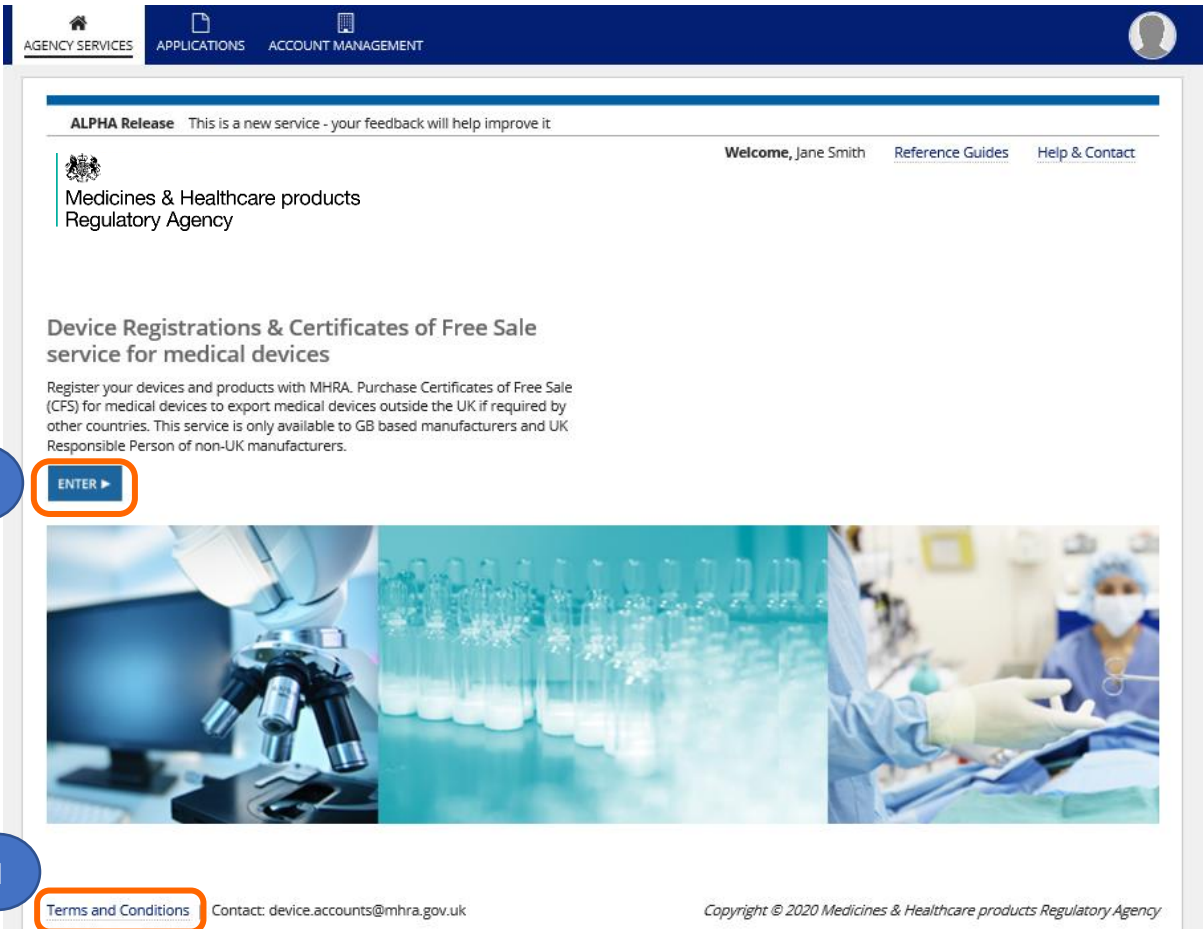
Since 01 January 2021 new medical device regulations have applied in the UK. Depending on your location and the type of Conformity Assessment that your registered device(s) comply with, the CFS certificate will state that the devices can be placed on either the GB (England, Scotland, Wales) market, or the UK (England, Scotland, Wales and Northern Ireland) market. For Northern Ireland-based manufacturers and Authorised Representatives with devices that are CE marked, the CFS certificate will also state that devices can be placed on the EU/EEA market. Please ensure that you download the preview and check with the recipient country **before** placing your order.

CFS for medical devices are issued as pdf documents with electronic signature. Hard copies with wet signature are no longer issued. We will try to process your order within 10 working days from date of receipt. During busy periods orders may take longer. Please order as far in advance as possible.

**Please note** that the Foreign, Commonwealth & Development Office will not apostille pdf documents directly from MHRA. You will need a Notary to carry out checks on the certificates with MHRA to ascertain their authenticity, and can then notarise the certificates, if required by the recipient country. Once notarised, the Foreign, Commonwealth & Development Office will apostille the documents.


**Where appropriate, Certificates of Free Sale are issued as a service to UK exporters. A Certificate of Free Sale should not be taken as a Government endorsement of any product that is referred to on the certificate. In issuing Certificates of Free Sale, the MHRA does not assess or verify that the product complies with relevant export requirements and restrictions. It is the applicant's responsibility to ensure compliance with these requirements and restrictions.**

1. **Read** our [terms and conditions](#). Certificate of Free Sale orders are non-refundable.
2. **Click** the [Enter](#) button on the [Landing](#) page.



**ALPHA Release** This is a new service - your feedback will help improve it

Welcome, Jane Smith [Reference Guides](#) [Help & Contact](#)

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### Device Registrations & Certificates of Free Sale service for medical devices

Register your devices and products with MHRA. Purchase Certificates of Free Sale (CFS) for medical devices to export medical devices outside the UK if required by other countries. This service is only available to GB based manufacturers and UK Responsible Person of non-UK manufacturers.

**2** [ENTER >](#)

**1** [Terms and Conditions](#) Contact: [device.accounts@mhra.gov.uk](mailto:device.accounts@mhra.gov.uk)

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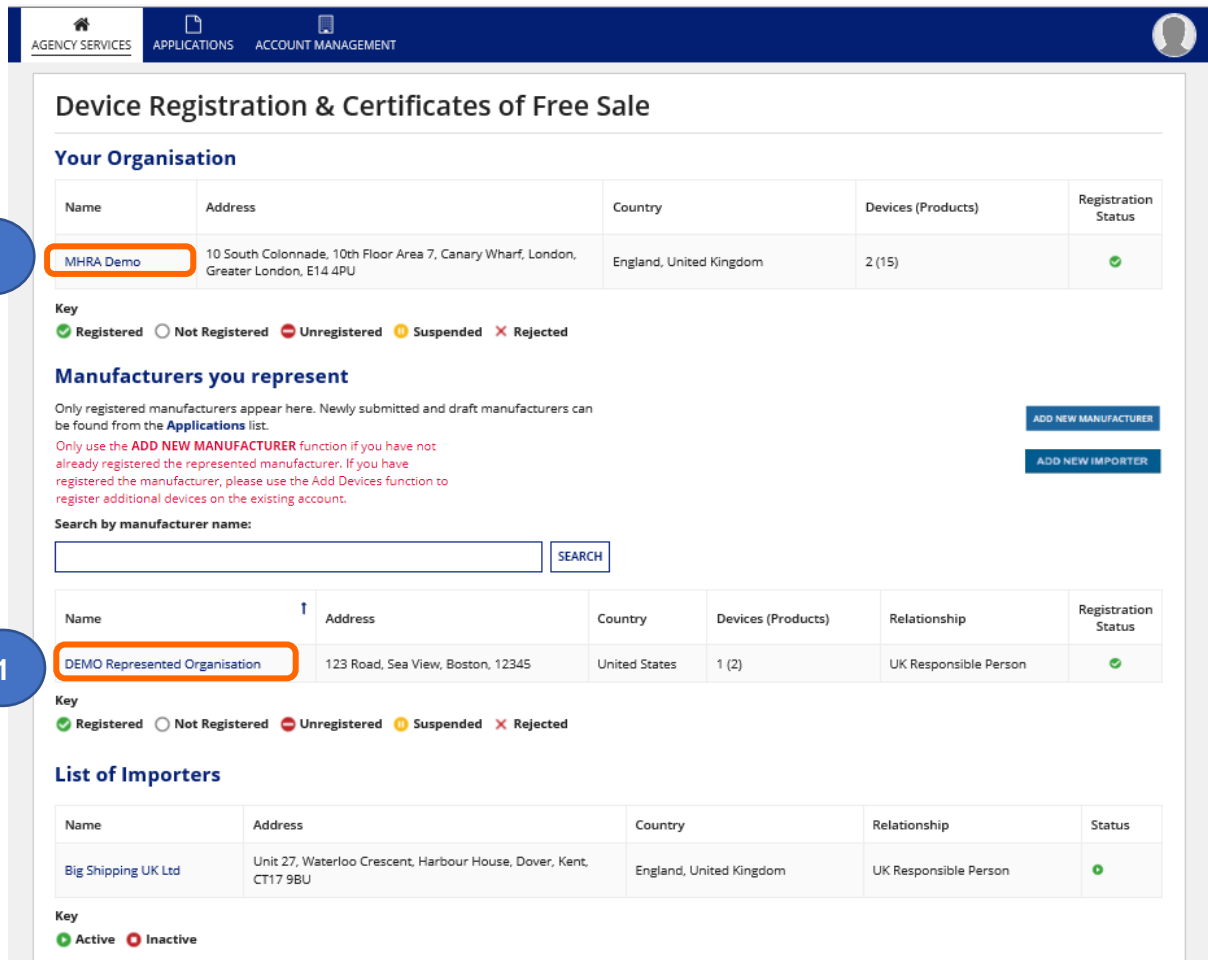


# Organisations

## Organisation page

1. **Click** on the **name** to select the **manufacturer** that you want to order a CFS for. This could be **Your Organisation** if you are a manufacturer or a **Represented Manufacturer** if you are a UK Responsible Person in the UK or an Authorised Representative in Northern Ireland.

**Please note** you can only order Certificates of Free Sale for **registered** devices. Follow all the instructions in the **Device Registration Reference Guide** to **Add devices** for your organisation or to **Add manufacturers** that you represent.



**Device Registration & Certificates of Free Sale**

**Your Organisation**

Name	Address	Country	Devices (Products)	Registration Status
MHRA Demo	10 South Colonnade, 10th Floor Area 7, Canary Wharf, London, Greater London, E14 4PU	England, United Kingdom	2 (15)	Registered

Key  
 Registered (green check) Not Registered (grey circle) Unregistered (red minus) Suspended (yellow exclamation) Rejected (red X)

**Manufacturers you represent**

Only registered manufacturers appear here. Newly submitted and draft manufacturers can be found from the **Applications** list.

Only use the **ADD NEW MANUFACTURER** function if you have not already registered the represented manufacturer. If you have registered the manufacturer, please use the Add Devices function to register additional devices on the existing account.

Search by manufacturer name:

Name	Address	Country	Devices (Products)	Relationship	Registration Status
DEMO Represented Organisation	123 Road, Sea View, Boston, 12345	United States	1 (2)	UK Responsible Person	Registered

Key  
 Registered (green check) Not Registered (grey circle) Unregistered (red minus) Suspended (yellow exclamation) Rejected (red X)

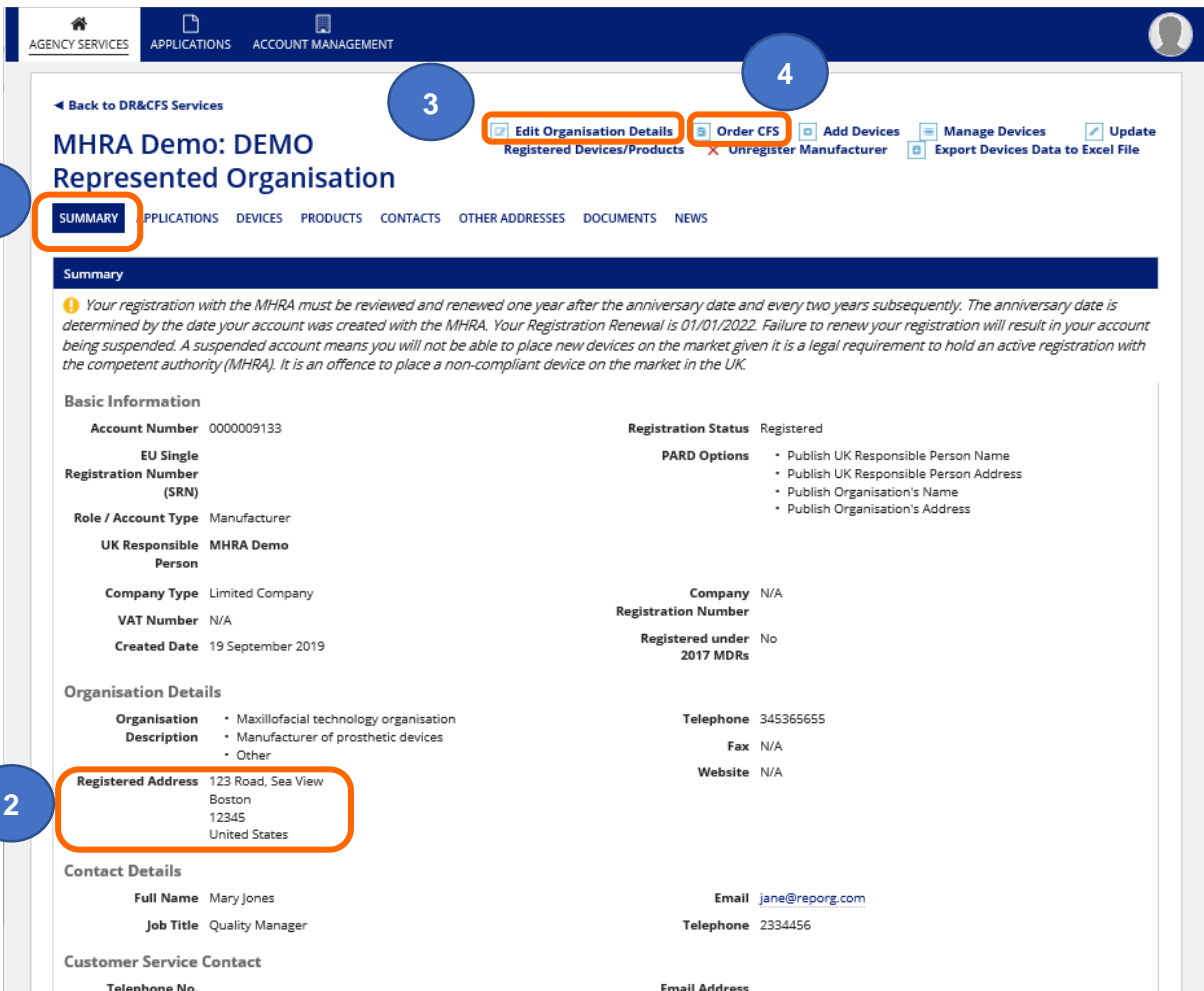
**List of Importers**

Name	Address	Country	Relationship	Status
Big Shipping UK Ltd	Unit 27, Waterloo Crescent, Harbour House, Dover, Kent, CT17 9BU	England, United Kingdom	UK Responsible Person	Active

Key  
 Active (green dot) Inactive (red dot)

## Check organisation information

1. **Check** that the organisation information is correct on the **Summary** page.
2. The **Registered Address** displayed here will be the address printed as the **Legal Manufacturer** on the Certificate of Free Sale.
3. If you need to edit organisation details, see the **Editing Organisation Details** section of the **Account Management Reference Guide**. **Statutory fees** apply to edit organisation details.
4. If you have already registered devices you will see the option to **Order CFS**. If no devices and products have been added, follow all the instructions in the **Device Registration Reference Guide** to **Add devices** for your organisation or a manufacturer that you represent as a UK Responsible Person in the UK or an Authorised Representative in Northern Ireland.



The screenshot shows the 'Summary' page for a 'MHRA Demo: DEMO Represented Organisation'. The page is annotated with four callouts:

- 1**: Points to the 'SUMMARY' tab in the navigation menu.
- 2**: Points to the 'Registered Address' field, which contains: 123 Road, Sea View, Boston, 12345, United States.
- 3**: Points to the 'Edit Organisation Details' button.
- 4**: Points to the 'Order CFS' button.

The page content includes a warning about registration renewal, a 'Basic Information' section with fields like Account Number, Registration Number, and Role, and an 'Organisation Details' section with a description and contact information.

Basic Information	
Account Number	0000009133
EU Single Registration Number (SRN)	
Role / Account Type	Manufacturer
UK Responsible Person	MHRA Demo Person
Company Type	Limited Company
VAT Number	N/A
Created Date	19 September 2019
Registration Status	Registered
PARD Options	<ul style="list-style-type: none"> <li>Publish UK Responsible Person Name</li> <li>Publish UK Responsible Person Address</li> <li>Publish Organisation's Name</li> <li>Publish Organisation's Address</li> </ul>
Company Registration Number	N/A
Registered under 2017 MDRs	No

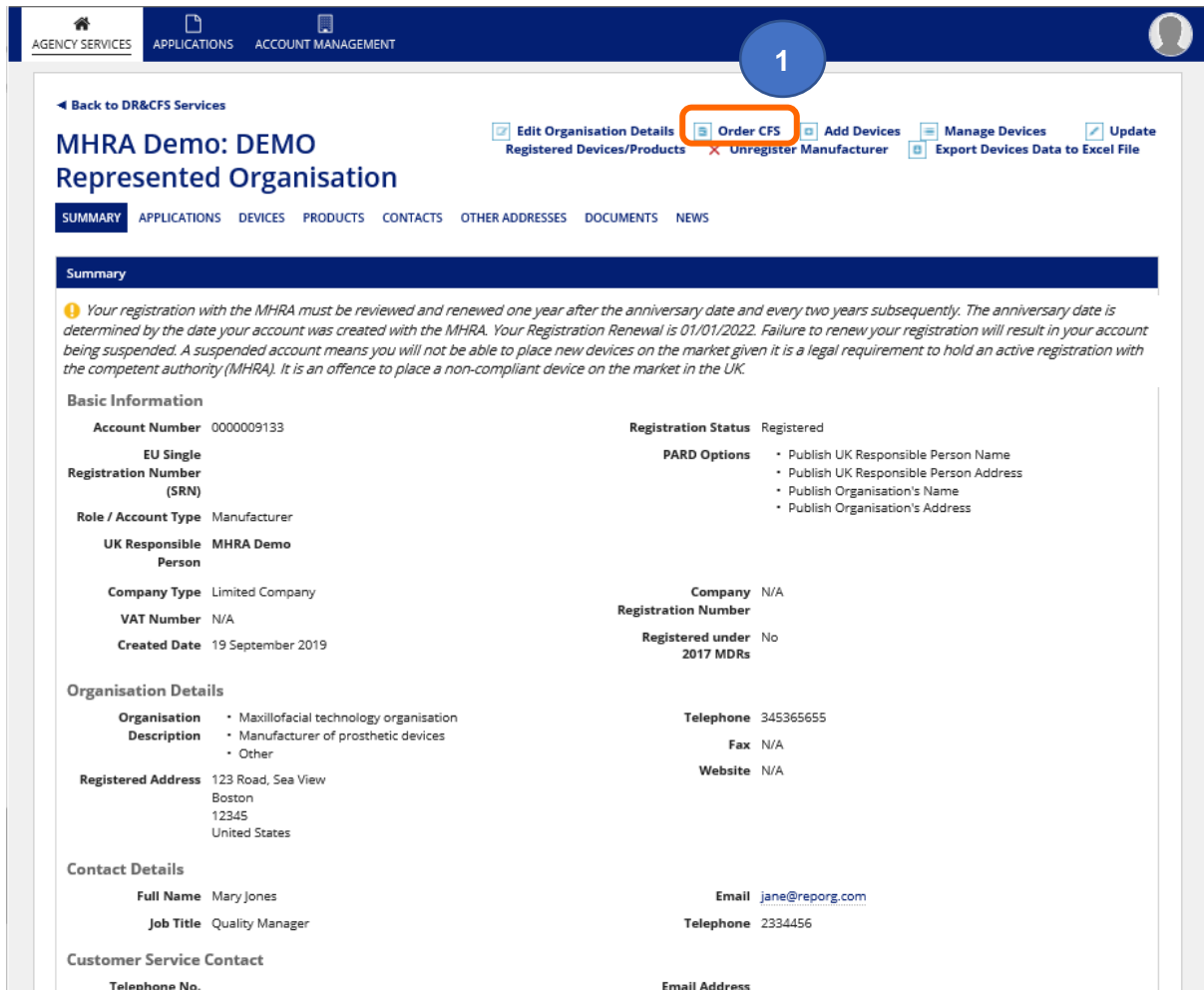
Organisation Details	
Organisation Description	<ul style="list-style-type: none"> <li>Maxillofacial technology organisation</li> <li>Manufacturer of prosthetic devices</li> <li>Other</li> </ul>
Telephone	345365655
Fax	N/A
Website	N/A

Contact Details	
Full Name	Mary Jones
Job Title	Quality Manager
Email	jane@reporg.com
Telephone	2334456

Customer Service Contact	
Telephone No.	
Email Address	

# Ordering a CFS

1. Click the Order CFS button.



The screenshot shows the MHRA Demo web application interface. At the top, there is a navigation bar with 'AGENCY SERVICES', 'APPLICATIONS', and 'ACCOUNT MANAGEMENT'. A blue circle with the number '1' is positioned above the 'Order CFS' button in the top right area of the page. The main content area is titled 'MHRA Demo: DEMO Represented Organisation'. Below the title, there are several action buttons: 'Edit Organisation Details', 'Order CFS' (highlighted with a red circle), 'Add Devices', 'Manage Devices', and 'Update Registered Devices/Products'. There is also a red 'X' icon next to 'Unregister Manufacturer' and a button for 'Export Devices Data to Excel File'. A 'Summary' section is visible, containing a warning message about registration renewal and a table of 'Basic Information' and 'Organisation Details'.

**Basic Information**

<b>Account Number</b>	0000009133	<b>Registration Status</b>	Registered
<b>EU Single Registration Number (SRN)</b>		<b>PARD Options</b>	<ul style="list-style-type: none"> <li>• Publish UK Responsible Person Name</li> <li>• Publish UK Responsible Person Address</li> <li>• Publish Organisation's Name</li> <li>• Publish Organisation's Address</li> </ul>
<b>Role / Account Type</b>	Manufacturer	<b>Company</b>	N/A
<b>UK Responsible Person</b>	MHRA Demo Person	<b>Registration Number</b>	
<b>Company Type</b>	Limited Company	<b>Registered under 2017 MDRs</b>	No
<b>VAT Number</b>	N/A		
<b>Created Date</b>	19 September 2019		

**Organisation Details**

<b>Organisation Description</b>	<ul style="list-style-type: none"> <li>• Maxillofacial technology organisation</li> <li>• Manufacturer of prosthetic devices</li> <li>• Other</li> </ul>	<b>Telephone</b>	345365655
<b>Registered Address</b>	123 Road, Sea View Boston 12345 United States	<b>Fax</b>	N/A
		<b>Website</b>	N/A

**Contact Details**

<b>Full Name</b>	Mary Jones	<b>Email</b>	jane@reporg.com
<b>Job Title</b>	Quality Manager	<b>Telephone</b>	2334456

**Customer Service Contact**

<b>Telephone No.</b>		<b>Email Address</b>	
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## Completing the CFS Application

### Adding products to your CFS order

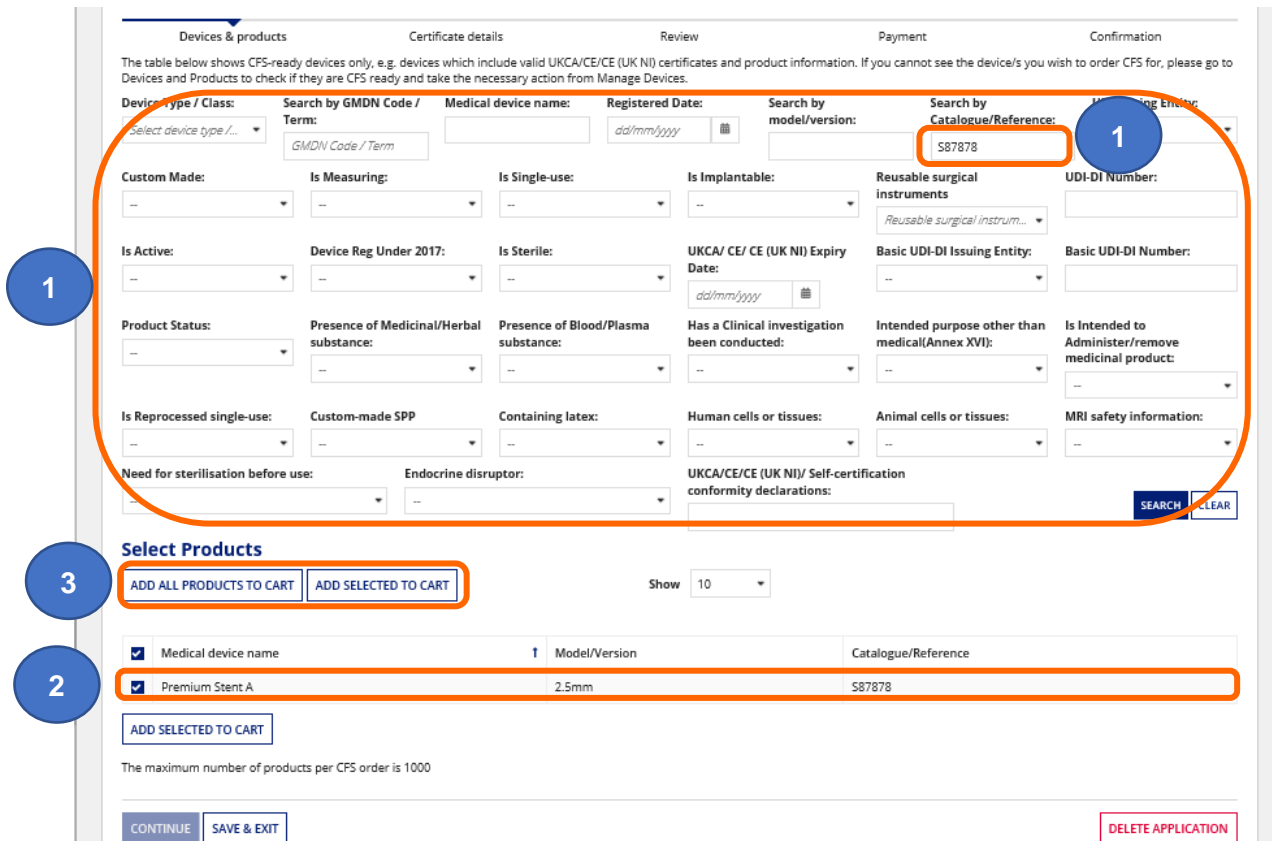
1. Use the **filters** to **search** for specific products. Only products that are **CFS Ready** will display, *in vitro diagnostic* medical devices for performance evaluation cannot be included on CFS orders so will not appear in the product table. If the conformity assessment document linked to the devices has expired, you cannot order CFS for the underlying products.

**Please note** if products you are expecting to see do not appear, check if conformity assessment document has expired - see [Manage registered devices](#) in the **Device Registration Reference Guide**.

If product information is incorrect see [Updating registered devices and products](#) in the **Device Registration Reference Guide**. You may need to remove the product and add it again.

2. **Select** the **products** which are to appear on the Certificate of Free Sale. **A maximum of 1000 products can be added to the order.**
3. Use the **Add to Cart** function to add your selected products. If you have 1000 products or less, you can **Add all products to cart**. If you select more than one product, your products will appear on the **Schedule** attached to the CFS Certificate.

**Please note** it is the product data that is added to the CFS order (**Medical Device Name, Model/Version** and **Catalogue/Reference only**) not the device (GMDN® Code or Term).



The screenshot displays the 'Devices & products' section of the CFS application. It includes a search form with various filters and a table of products. Three red annotations highlight key features:

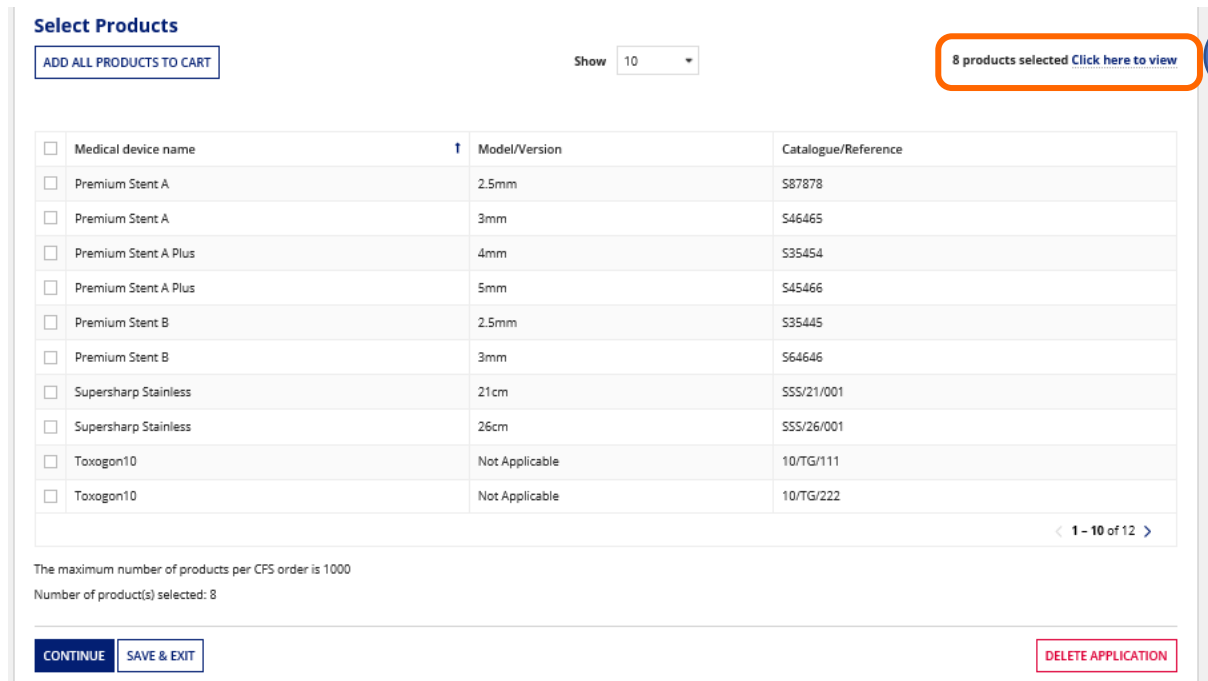
- 1:** A red circle highlights the search filters and input fields, including 'Search by Catalogue/Reference' with the value 'S87878'.
- 2:** A red circle highlights a row in the product table, which is also enclosed in a red box. The row contains:
 

<input checked="" type="checkbox"/>	Medical device name	1	Model/Version	Catalogue/Reference
<input checked="" type="checkbox"/>	Premium Stent A		2.5mm	S87878
- 3:** A red circle highlights the 'ADD ALL PRODUCTS TO CART' and 'ADD SELECTED TO CART' buttons.

At the bottom of the interface, there are buttons for 'CONTINUE', 'SAVE & EXIT', and 'DELETE APPLICATION'. A note states: 'The maximum number of products per CFS order is 1000'.

## Check products in cart

- When you have added all the required products to cart [click](#) to [view cart](#).



**Select Products**

[ADD ALL PRODUCTS TO CART](#) Show 10 8 products selected [Click here to view](#)

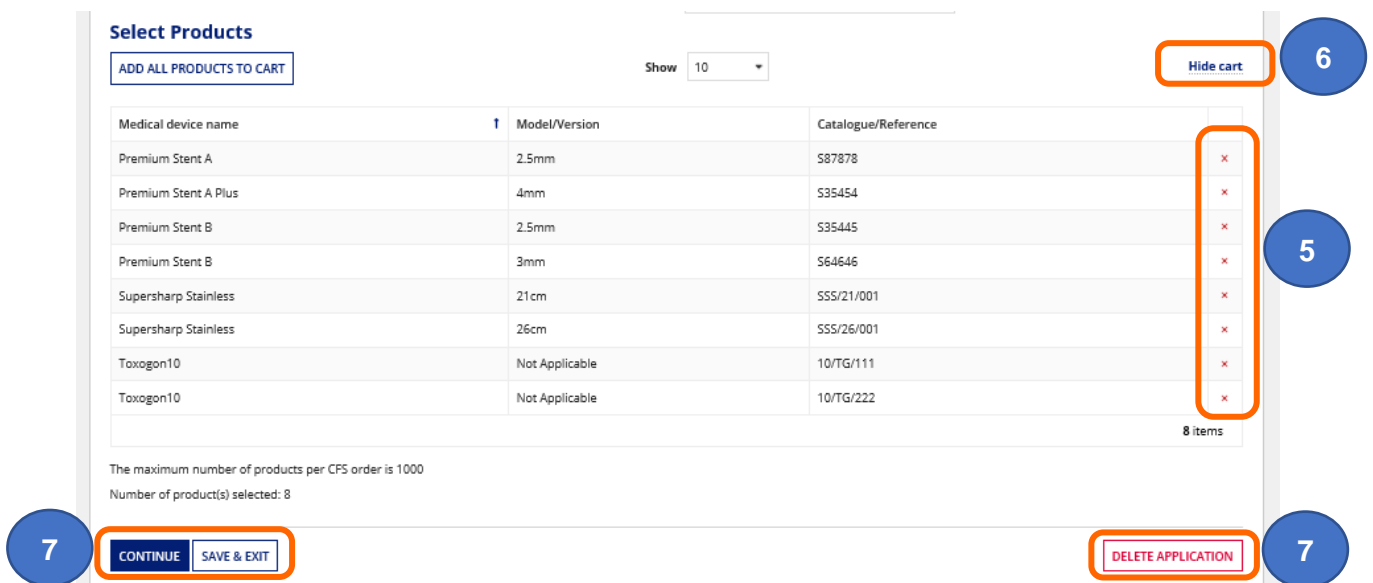
<input type="checkbox"/>	Medical device name	Model/Version	Catalogue/Reference
<input type="checkbox"/>	Premium Stent A	2.5mm	S87878
<input type="checkbox"/>	Premium Stent A	3mm	S46465
<input type="checkbox"/>	Premium Stent A Plus	4mm	S35454
<input type="checkbox"/>	Premium Stent A Plus	5mm	S45466
<input type="checkbox"/>	Premium Stent B	2.5mm	S35445
<input type="checkbox"/>	Premium Stent B	3mm	S64646
<input type="checkbox"/>	Supersharp Stainless	21cm	SSS/21/001
<input type="checkbox"/>	Supersharp Stainless	26cm	SSS/26/001
<input type="checkbox"/>	Toxogon10	Not Applicable	10/TG/111
<input type="checkbox"/>	Toxogon10	Not Applicable	10/TG/222

< 1 - 10 of 12 >

The maximum number of products per CFS order is 1000  
Number of product(s) selected: 8

[CONTINUE](#) [SAVE & EXIT](#) DELETE APPLICATION

- The products added will display. If you want to remove any [click](#) the red **X** to remove.



**Select Products**

[ADD ALL PRODUCTS TO CART](#) Show 10 Hide cart

<input type="checkbox"/>	Medical device name	Model/Version	Catalogue/Reference	
<input type="checkbox"/>	Premium Stent A	2.5mm	S87878	X
<input type="checkbox"/>	Premium Stent A Plus	4mm	S35454	X
<input type="checkbox"/>	Premium Stent B	2.5mm	S35445	X
<input type="checkbox"/>	Premium Stent B	3mm	S64646	X
<input type="checkbox"/>	Supersharp Stainless	21cm	SSS/21/001	X
<input type="checkbox"/>	Supersharp Stainless	26cm	SSS/26/001	X
<input type="checkbox"/>	Toxogon10	Not Applicable	10/TG/111	X
<input type="checkbox"/>	Toxogon10	Not Applicable	10/TG/222	X

8 items

The maximum number of products per CFS order is 1000  
Number of product(s) selected: 8

CONTINUE SAVE & EXIT DELETE APPLICATION

- [Click](#) the [Hide cart](#) link to add more products. If adding more products, ensure you view cart again before proceeding.
- [Click](#) the [Continue](#) button to continue to Review page or [Save & Exit](#) to save a TEMP application or [Delete Application](#) to discard or start again.

## Certificate details

1. You can have **one** address in addition to the legal manufacturer's address on the CFS certificate. You can select which additional address appears on the Certificate.
2. You can select more addresses to appear on the [schedule](#). Both UKRP and Authorised Representative addresses as well as manufacturer [other addresses](#) will appear in the dropdown. Ensure that you select an address that has been added for this specific manufacturer.

**Important note:** the [Clear More addresses](#) button is **not** currently working. If you select incorrect addresses you will need to **delete** the application and start again. We are working towards a fix for this issue. Please accept our apologies for any inconvenience and ensure that you [Review order and preview CFS](#) before making payment.

**Address to be printed on CFS**

Registered Address: MANAGE ADDRESSES  
10 South Colonnade, 10th Floor, Cabot Square, Canary Wharf, London, E14  
4PU, England, United Kingdom

One additional address on certificates:  
[Manufacturer] [Manufacturing site/Physical manufacturer] China Medical Co, No 7, Section 5, Luzhongshan, Liaoning, 110001, China

You can choose only one additional address to be printed on certificates.

**More addresses on schedule:**  
[Manufacturer] [Manufacturing site/Physical manufacturer] India Medical Co, Andhra Cantt, Hyderabad, Andhra Pradesh, 523270, India

Any other addresses you choose will be printed on the schedule.


**CLEAR MORE ADDRESSES**

3. You can add additional addresses to the system by clicking on the [Manage addresses](#) link. Follow the steps in the [Shipping, Billing, Manufacturing Site Addresses](#) section of the MHRA **Account Management Reference Guide** to add addresses.

**Please note** if you click on the [Manage addresses](#) link you will be asked to [save](#) your application. You can then proceed to Manage addresses then come back and complete your application by following the [Save and exit: resume applications](#) instructions.

4. You no longer need to select a [Certificate delivery address](#) as all CFS orders are now issued as pdf documents with electronic signature. Hard copies with wet signature are no longer issued. The pdf will be emailed to the email address of the person who placed the order.

AGENCY SERVICES    APPLICATIONS    ACCOUNT MANAGEMENT
 



---

## CFS (Certificate of Free Sale) Application for DEMO Represented Organisation - TEMP20220221162325

Devices & products
Certificate details
Review
Payment
Confirmation

### Address to be printed on CFS

**Registered Address:** MANAGE ADDRESSES 3

123 Road, Sea View, Boston, 12345, United States

**One additional address on certificates:** 1

[UK Responsible Person] [Billing Address] MHRA Finance Dept, 10, South Colonnade, LONDON, London, E14 4PU, United Kingdom

You can choose only one additional address to be printed on certificates.

**More addresses on schedule:** 2

[Manufacturer] [Manufacturing site/Physical manufacturer] Mexico Medical inc, 678 Buenavista , Mexico City, Iztacalo, 00810, Mexico

Any other addresses you choose will be printed on the schedule.

### Special formatting request (optional)

The only request that will be considered is for manufacturing site address/s where no schedule is generated due to only one product (model/version) on the order. You must use the 'Other Addresses' function to add manufacturing site addresses for selection on CFS orders where more than one product is added to the order and a schedule will be generated. Only the Medical Device Name, Model/Version and Catalogue/Reference columns will be included on CFS orders. All orders are sorted by Medical Device Name. Please do not ask for other formatting changes as they will not be considered and CFS orders are non-refundable.

5

### Certificate delivery address

Registered address: MANAGE ADDRESSES 3

10 South Colonnade, 10th Floor Area 7, Canary Wharf, London, Greater London, E14 4PU, England, United Kingdom

Alternative address:

4

5. Enter any [Special Formatting Requests](#).

**Please note** MHRA will not:

- add other addresses to the Certificate or Schedule **unless** the order is for a **single** product (where no schedule is generated) – you need to add and select other addresses **before** making payment. Follow the steps in the [Shipping, Billing, Manufacturing Site Addresses](#) section of the MHRA **Account Management Reference Guide** to add other addresses
- make **any** changes to CFS certificate or schedule including layout or text or adding additional columns

6. Enter the [country name/s](#) and [number of certificates](#). If you do not want to specify a country, tick the 'Do not specify a country' box. Check that the recipient country will accept this.

**Please note:**

You can only select from countries that appear in the [Foreign, Commonwealth & Development Office's \(FCDO\) Geographical names index](#). We print the Country Name from this index on the CFS certificate e.g. for CFS requested for Venezuela we print 'Venezuela', not the Official Name e.g. 'The Bolivarian Republic of Venezuela'.

If the country name does not appear in the dropdown in the system, or in the above index, you can still order a CFS by selecting 'Do not specify a country'. However, you must check that the recipient country will accept this, before placing your order.

The [fee](#) is dependant on the total number of certificates.

The expiry date on all CFS certificates will be the earliest Conformity Assessment document expiry date of any device included in the CFS order plus 365 days. For devices with Declaration of Conformity or Custom-made Statements the maximum validity will be 5 years.

7. Click the [Continue](#) button to review your order.

**Enter countries and number of certificates**

Country	Number of certificates		
Ghana ✕	2	<input type="checkbox"/> Do not specify a country	✕
United Arab Emirates ✕	2	<input type="checkbox"/> Do not specify a country	✕
<a href="#">+ Add country</a>			

Total number of certificates: 4  
 Price: £75.00

We charge £75.00 for 1 to 10 certificates and £10.00 for each additional certificate

[CONTINUE](#)
[SAVE & EXIT](#)
[BACK](#)
[DELETE APPLICATION](#)



## Review order and preview CFS Certificate and Schedule

1. **Review** and/or **Edit** your Order.
2. **Click** the **Download CFS documents Preview** link to view and review the CFS certificate and schedule for your order.

**Please note:** Our system allows you to **preview** an order **before** submitting payment. It is your responsibility to ensure that the order is correct and acceptable to the receiving country. **CFS orders are non-refundable** – see our terms and conditions.

If you forget to preview your schedule and certificate before you press the **Continue to Payment button**, and then click the **Back** button on the payment page, the previews will no longer be available. To preview you will need to go **Back** again to **Order page** and then **click** the **continue** button. The preview will then be available again.

3. If you need to make any changes **click** the **Edit** button.

**Important note:** the **Clear More addresses** button is **not** currently working. If you select incorrect addresses you will need to **delete** the application and start again. We are working towards a fix for this issue. Please accept our apologies for any inconvenience and ensure that you **Review order and preview CFS** before making payment.

**Address to be printed on CFS**

Registered Address: MANAGE ADDRESSES  
10 South Colonnade, 10th Floor, Cabot Square, Canary Wharf, London, E14  
4PU, England, United Kingdom

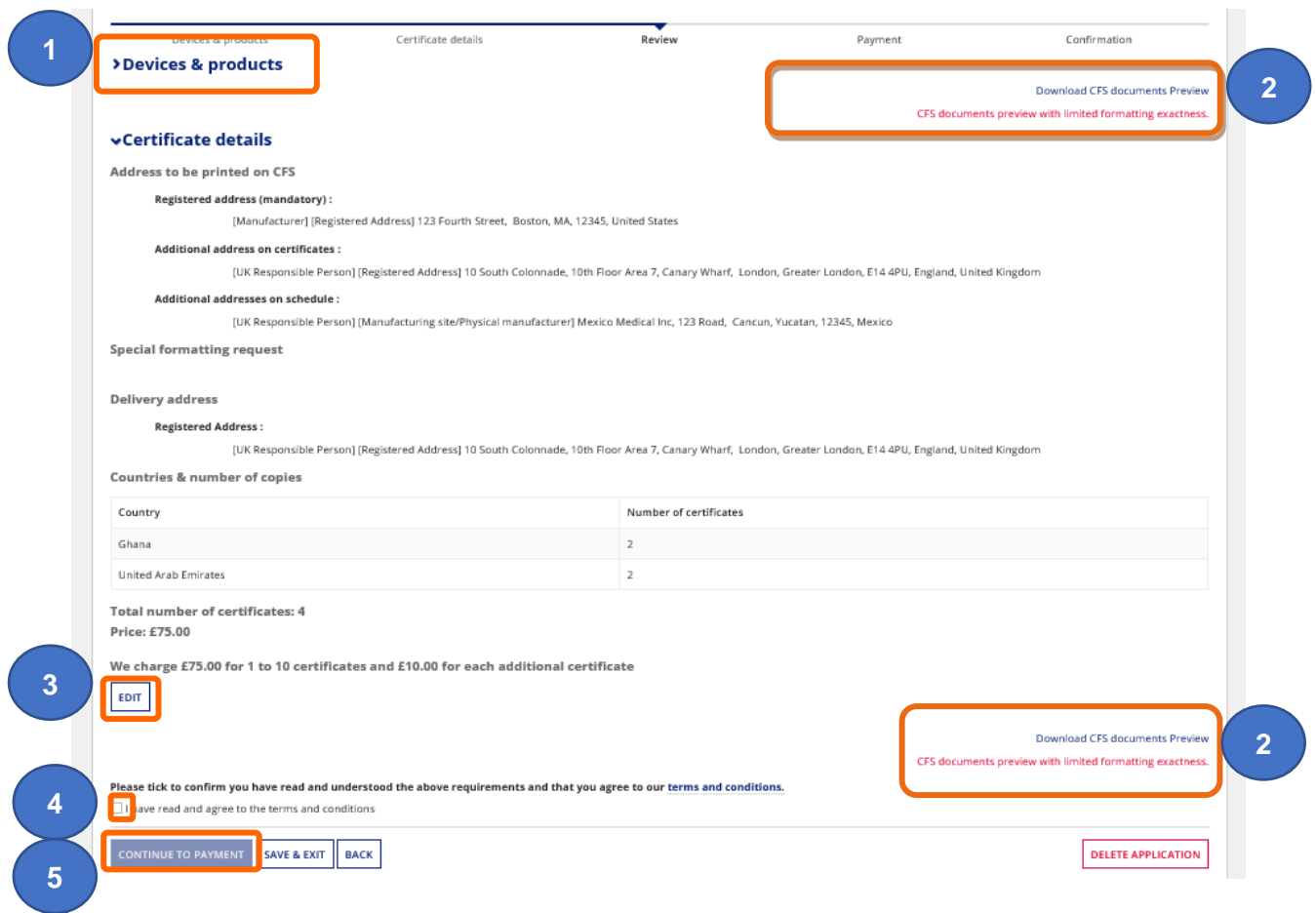
One additional address on certificates:  
[Manufacturer] [Manufacturing site/Physical manufacturer] China Medical Co, No 7, Section 5, Luzhongshan, Liaoning, 110001, China

You can choose only one additional address to be printed on certificates.

**More addresses on schedule:**  
[Manufacturer] [Manufacturing site/Physical manufacturer] India Medical Co, Andhra Cantt, Hyderabad, Andhra Pradesh, 523270, India

Any other addresses you choose will be printed on the schedule.

**CLEAR MORE ADDRESSES**



The screenshot shows the 'Review' stage of a CFS application. The breadcrumb trail includes 'Devices & products', 'Certificate details', 'Review', 'Payment', and 'Confirmation'. Callout 1 points to the 'Devices & products' breadcrumb. Callout 2 points to a 'Download CFS documents Preview' button with a warning: 'CFS documents preview with limited formatting exactness.' Callout 3 points to an 'EDIT' button. Callout 4 points to a checkbox labeled 'I have read and agree to the terms and conditions'. Callout 5 points to the 'CONTINUE TO PAYMENT' button. Other buttons visible are 'SAVE & EXIT', 'BACK', and 'DELETE APPLICATION'.

4. **Read & Tick** the ‘I have read and agree to the terms and conditions’ checkbox if you have read and agree to the terms and conditions.

5. **Click** the **Continue to Payment** button.

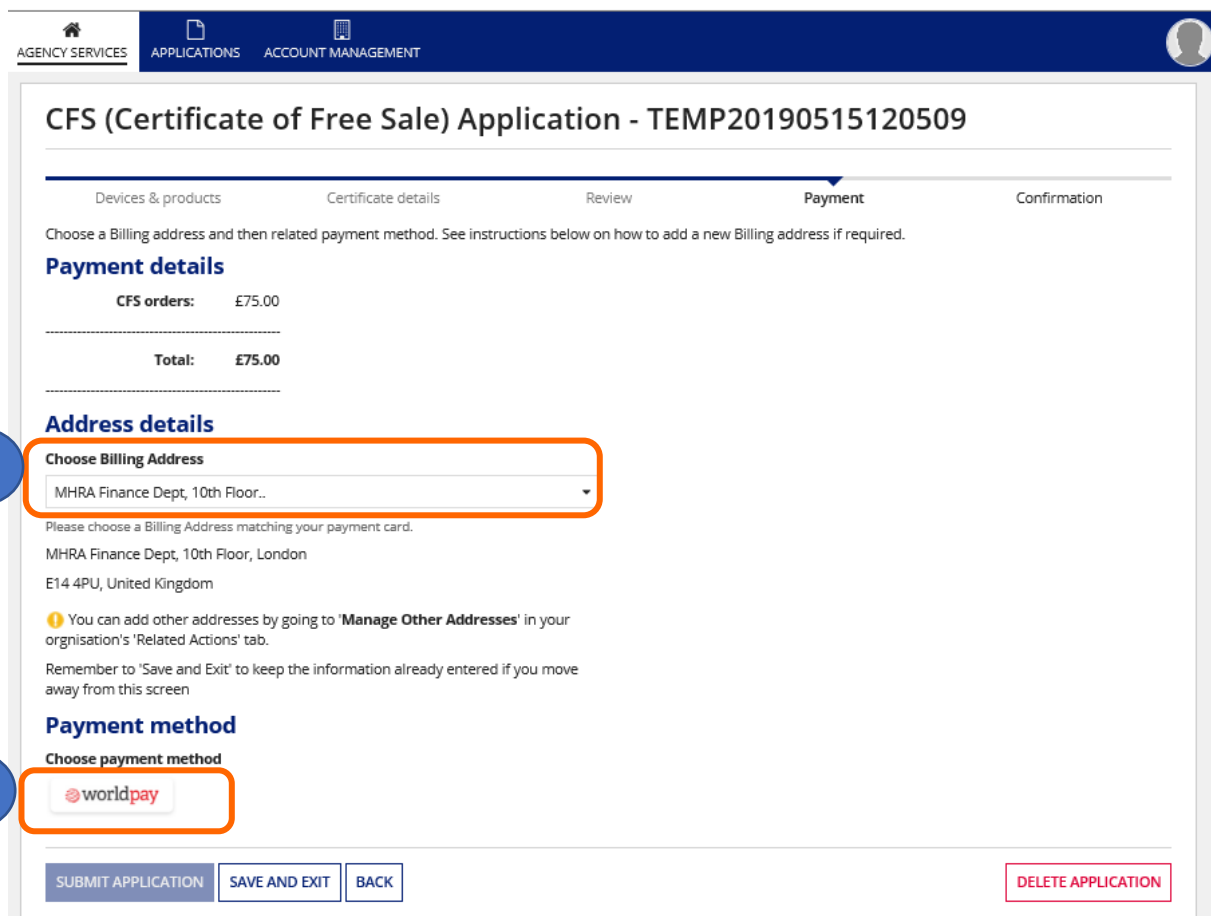
## Pay for your CFS order

### 1. Chose a billing address

See [Managing Shipping, Billing, Manufacturing Site addresses](#) in the **MHRA Account Management Reference Guide**.

### 2. Click on the [worldpay](#) button. We only accept payment by worldpay for CFS orders.

**Important note:** Fee in screenshot is for illustrative purposes only. Check current [fee](#) on our website.



**CFS (Certificate of Free Sale) Application - TEMP20190515120509**

Devices & products   Certificate details   Review   **Payment**   Confirmation

Choose a Billing address and then related payment method. See instructions below on how to add a new Billing address if required.

**Payment details**

CFS orders: £75.00

Total: £75.00

**Address details**

1 Choose Billing Address

MHRA Finance Dept, 10th Floor..

Please choose a Billing Address matching your payment card.

MHRA Finance Dept, 10th Floor, London  
E14 4PU, United Kingdom

**!** You can add other addresses by going to 'Manage Other Addresses' in your organisation's 'Related Actions' tab.

Remember to 'Save and Exit' to keep the information already entered if you move away from this screen

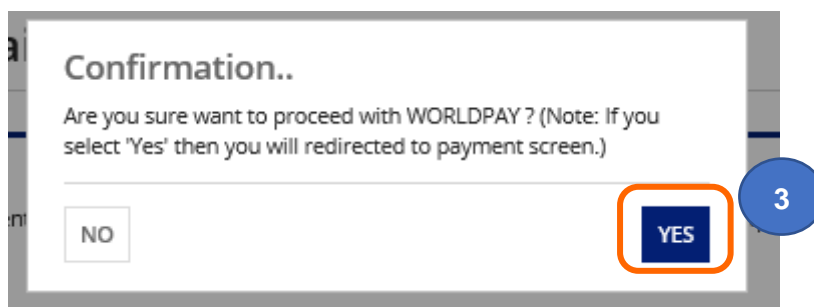
**Payment method**

2 Choose payment method

worldpay

SUBMIT APPLICATION   SAVE AND EXIT   BACK   DELETE APPLICATION

### 3. A confirmation message will appear. Select the Yes button if you wish to proceed

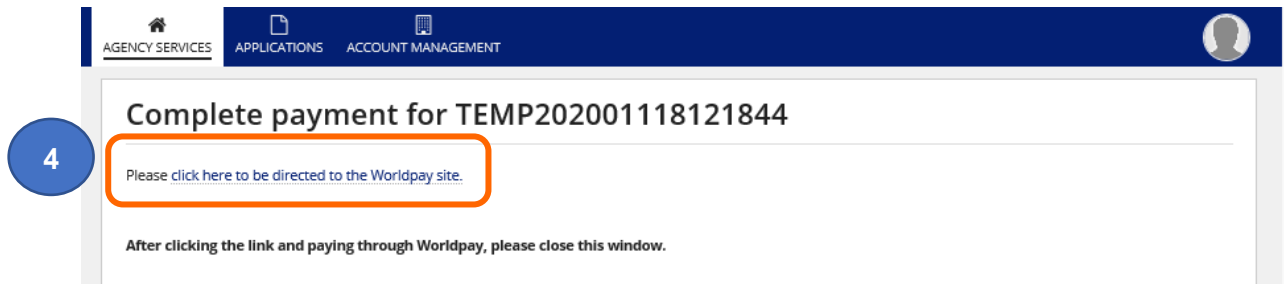


**Confirmation..**

Are you sure want to proceed with WORLDPAY ? (Note: If you select 'Yes' then you will redirected to payment screen.)

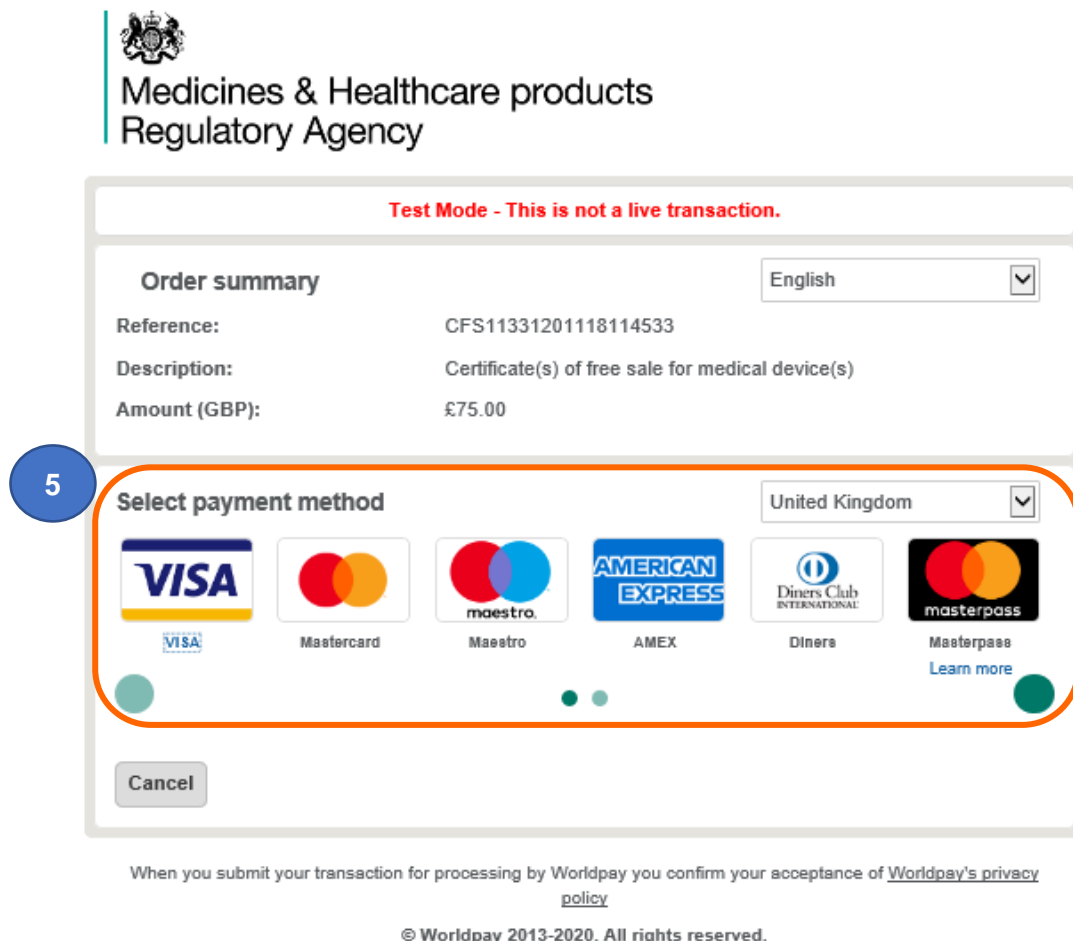
NO   YES

- Click the link to be directed to the [worldpay](#) site.



- Select the [payment method](#).

**Important note:** Fee in screenshot is for illustrative purposes only. Check current [fee](#) on our website.



- Enter payment details and [click](#) the [Make payment](#) button.

**Important note:** Fee in screenshot is for illustrative purposes only. Check current [fee](#) on our website.

**Test Mode - This is not a live transaction.**

---



**Order summary** English

Reference: CFS11331201118114533

Description: Certificate(s) of free sale for medical device(s)

Amount (GBP): £75.00

---


**Payment details**  

\* Indicates a required field

Card number \*

Cardholder's name \*

Expiry date \*  /


Security code \*   Last 3 digits on the back of card

---

**Billing address**  
DEMO Finance Department, 20 City Road, London , EC1 6ZH, United Kingdom

**Contact details**  
devices.transformation@mhra.gov.uk

- [Click](#) the [Submit Application](#) button. If you do not click this button (and your payment was successful) the TEMP application will remain in the [Applications Tab](#) and you will need to wait at least **24 hours** for the application to be auto-submitted to MHRA. Please ensure that you click [Submit Application](#) to avoid unnecessary issues and delays.

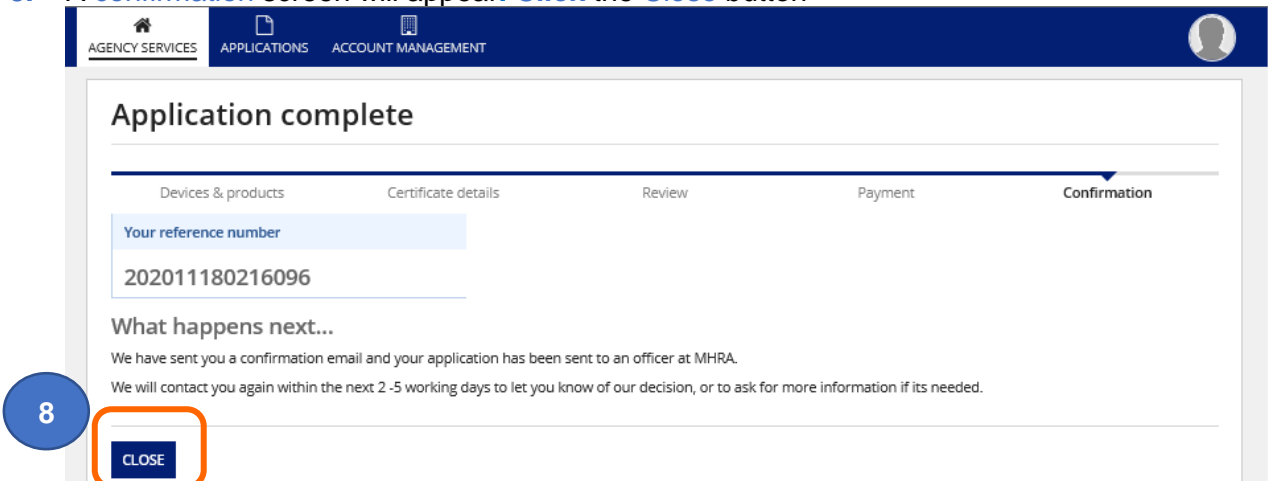
AGENCY SERVICES APPLICATIONS ACCOUNT MANAGEMENT 

---

**Payment complete**

Your payment completed successfully. Please submit your application

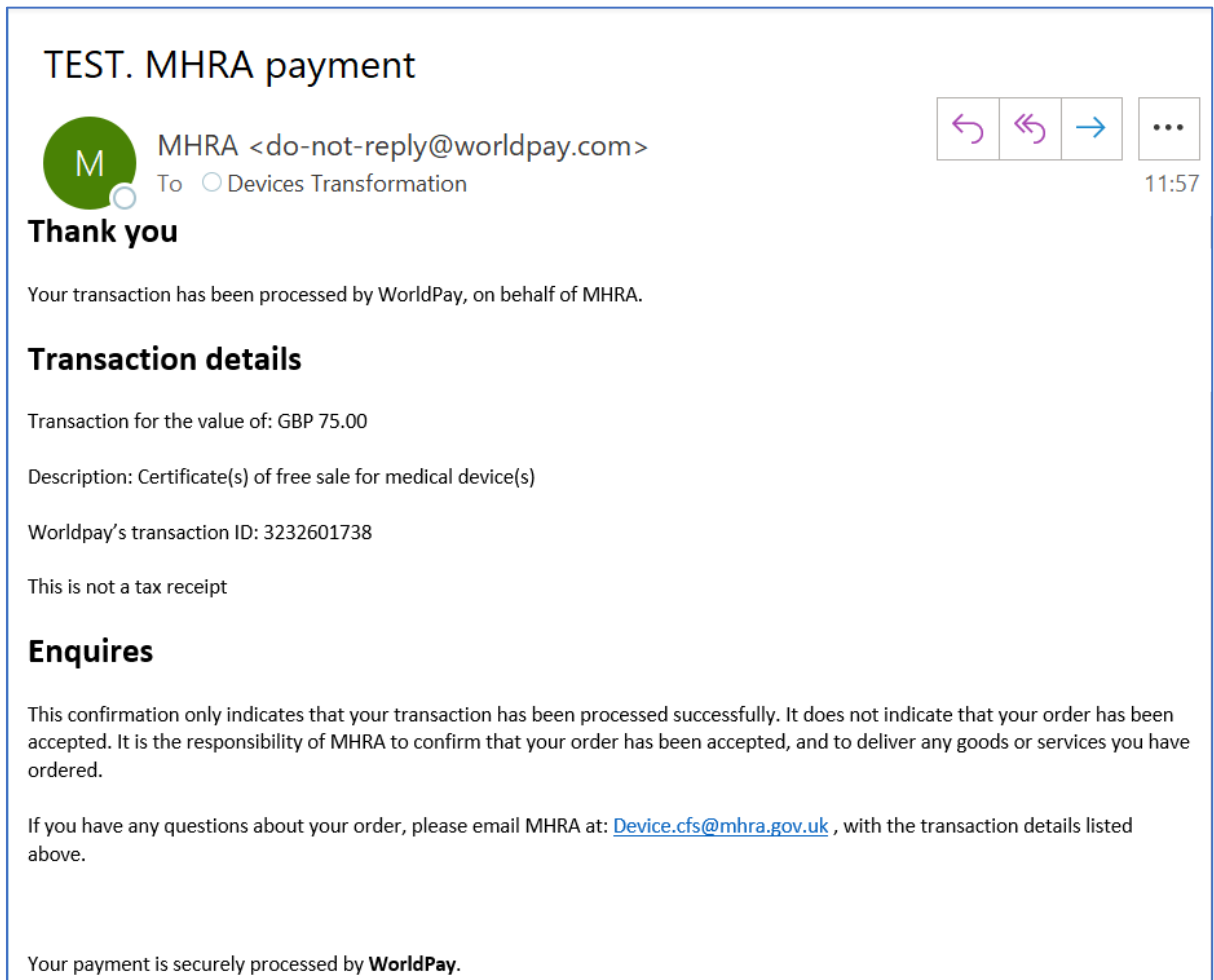
8. A confirmation screen will appear. Click the Close button



9. You will receive a confirmation email from worldpay.

**Please note** MHRA does not issue tax receipts. The worldpay transaction email and the Certificates of Free Sale order email and receipt are the only documents you will receive in relation to payment for your Certificates of Free Sale order.

**Important note:** Fee in screenshot is for illustrative purposes only. Check current [fee](#) on our website.









## CFS Order confirmation email

10. You will receive a [confirmation](#) email from MHRA.

**Important note:** Fee in screenshot is for illustrative purposes only. Check current [fee](#) on our website.

Certificate of Free Sale Service - DEMO Represented Organisation ...

 No Reply TEST <no-reply@mhra.gov.uk>  
To:  Devices Transformation

12:03

Dear Jane Smith,

We've received your CFS Order application on 18 November 2020.

Application reference number: 202011180216096

Manufacturer name(s)  
DEMO Represented Organisation

Cost of application (£): 75

Country	Quantity
Ghana	2
United Arab Emirates	2

We will check the information you've given us and will send you an email within the next 5 days to let you know if your request has been accepted or rejected. If you haven't received a reply from us within 5 days please check your junk mail folder.

Access your [MHRA account](#).

Remember: do not share your account details and keep them safe.  
MHRA won't accept responsibility for any unauthorised access to your account.

Yours sincerely,

**Certificates of free sale service**  
**Devices division**  
**MHRA**  
**10 South Colonnade, Canary Wharf, London, E14 4PU**


## Order complete confirmation





11. Once we have processed and emailed the pdf documents for your order in a **separate email**, you will receive an automated **confirmation** email from MHRA.

**Please note** CFS for medical devices are issued as pdf documents with electronic signature. Hard copies with wet signature are no longer issued. We will try to process your order within 10 working days from date of receipt. During busy periods orders may take longer. Please order as far in advance as possible.


**Please note** that the Foreign, Commonwealth & Development Office will not apostille pdf documents directly from us. You will need a Notary who will carry out checks on the certificates with MHRA to ascertain their authenticity, and can then notarise the certificates, if required by the recipient country. Once notarised, the Foreign, Commonwealth & Development Office will apostille the documents.

MHRA Certificates of Free Sale service -DEMO Represented Organ...

 No Reply TEST <no-reply@mhra.gov.uk>  
To  Devices Transformation

12:39

 This message was sent with High importance.

Dear Jane Smith,

Application reference: 202011180216096

Further to your Certificate of Free Sale order on 18 November 2020 for:

Manufacturer organisation:  
DEMO Represented Organisation

Address:  
123 Street  
Sea View Industrial Estate  
Boston  
MA  
12345  
United States

The PDF documents for your CFS order have been emailed to the user who submitted the order. We no longer issue hard copies.

The PDF(s) do not contain the watermark and hologram found on hard copies. They do have electronic signatures of MHRA staff as registered with the Foreign and Commonwealth Office. We will assist where possible, and if necessary, to verify that these documents have been created by MHRA should any receiving country have queries. All such queries must be sent to [Device.cfs@mhra.gov.uk](mailto:Device.cfs@mhra.gov.uk)

The account number for your company/organisation is 0000005365. [Sign into your account.](#)

Remember: do not share your account details and keep them safe.  
MHRA won't accept responsibility for any unauthorised access to your account.

Please do not respond directly to this email address. The originating email account is not monitored.



## CFS layout

**Please note** the expiry date on all CFS certificates will be the earliest Conformity Assessment document expiry date of any device included in the CFS order plus 365 days. For devices with Declaration of Conformity or Custom-made Statements the maximum validity will be 5 years.

 Medicines & Healthcare products Regulatory Agency	<b>Example only</b>	10 South Colonnade Canary Wharf London E14 4PU United Kingdom  mhra.gov.uk
Certificate number: <b>2023051202218001/1</b>		
Order date: <b>12/05/2023</b>		
CFS Expiry date: <b>31/10/2024</b>		
<b>Medicines and Healthcare products Regulatory Agency</b>		
On behalf of the Secretary of State for Health and Social Care		
<b>CERTIFICATE OF FREE SALE FOR EXPORTATION OF MEDICAL PRODUCTS TO GHANA</b>		
<p>It is hereby certified that, on the basis of information provided, and subject to approval requirements for covid tests (see below), the products named below and detailed in the attached schedule (if applicable), which are manufactured by: Legal Manufacturer: <b>DEMO Represented Organisation</b>, 123 Road, Sea View, Boston, 12345, United States UK Responsible Person: <b>MHRA Demo</b>, 10 South Colonnade, 10th Floor Area 7, Canary Wharf, London, Greater London, E14 4PU, England, United Kingdom have been affixed with the relevant conformity marking(s) e.g. CE under EU MDD 93/42/EEC, CE under EU IVDD 98/79/EC, UKCA under UK MDR (2002/618) Part II and may be freely sold in Great Britain (England, Scotland, Wales). In the UK, certain covid test devices must be approved by the Secretary of State before they can be freely placed on the market or put into service. UK approved tests are listed at: <a href="https://www.gov.uk/government/publications/covid-19-test-validation-approved-products">https://www.gov.uk/government/publications/covid-19-test-validation-approved-products</a>.</p> <p>Please see attached schedule</p> <p>Where appropriate, Certificates of Free Sale are issued as a service to UK exporters. A Certificate of Free Sale should not be taken as a Government endorsement of any product that is referred to on the certificate. In issuing Certificates of Free Sale, the MHRA does not assess or verify that the product complies with relevant export requirements and restrictions. It is the applicant's responsibility to ensure compliance with these requirements and restrictions.</p> <p>Yours sincerely,</p>		



## Schedule layout

**Please note** only the **Medical Device Name**, **Model/Version** and **Catalogue/Reference**, **Basic UDI DI** and **UKCA or CE Certificate Reference No.** data will appear on the CFS certificate or schedule. If any of these fields have not been populated in the registrations system, 'Not applicable' or N/A' or will appear on the CFS certificate or schedule. To populate these fields, where applicable, follow the [Update products](#) instructions in the **Device Registration Reference Guide**.

If you select more than one product, your products will appear on the [Schedule](#) attached to the CFS Certificate. Please also note that it is the product that is added to the CFS order (**Medical Device Name**, **Model/Version** and **Catalogue/Reference**, **Basic UDI DI** and **UKCA or CE Certificate Reference No.** **only**) not the device (GMDN<sup>®</sup> Code or Term).

Certificate Reference: 2023051202218001/1  
Ordered on: 12/05/2023

Directive/Regulation: CE under EU MDD 93/42/EEC, CE under EU IVDD 98/79/EC, UKCA under UK MDR (2002/618) Part II,

Manufacturer Name: DEMO Represented Organisation

CFS Expiry Date: 31/10/2024

Addresses:

[UK Responsible Person] Manufacturing site/Physical manufacturer: India Medical Co , Andhra Cantt, Hyderabad, Andhra Pradesh, 523270, India

[UK Responsible Person] Manufacturing site/Physical manufacturer: Mexico Medical Inc, 123 Road, Cancun, Yucatan, 12345, Mexico

#	Medical Device Name	Model/Version	Catalogue/Reference	Basic UDI DI	Certificate Reference No.
1	Altmatsys	Altmatsys1	AMS/001/01	N/A	BSI_6533453
2	Custom retainer	Cusret1	Not Applicable	N/A	N/A
3	Custom retainer-child	Cusretchild1	Not Applicable	N/A	N/A
4	Premium Stent A	3mm	S46465	8788787656	67868768, 5746576567_EU MDR Art120 Extension
5	Premium Stent A	2.5mm	S87878	8788787656	67868768, 5746576567_EU MDR Art120 Extension
6	Premium Stent A Plus	5mm	S45466	8788787656	67868768, 5746576567_EU MDR Art120 Extension
7	Premium Stent A Plus	4mm	S35454	8788787656	67868768, 5746576567_EU MDR Art120 Extension
8	Premium Stent B	3mm	S64646	8788787656	67868768, 5746576567_EU MDR Art120 Extension
9	Premium Stent B	2.5mm	S35445	8788787656	67868768, 5746576567_EU MDR Art120 Extension
10	Supersharp Stainless	21cm	SSS/21/001	657668787889898	67868768
11	Supersharp Stainless	26cm	SSS/26/001	657668787889898	67868768
12	Toxogon10	Not Applicable	10/TG/444	076568548548	546576767
13	Toxogon10	Not Applicable	10/TG/333	076568548548	546576767

Page 2 of 3

## Manage registered devices

If products need to be added or removed from an existing registered device or Conformity Assessment documents certificates need to be uploaded, linked, or removed (unlinked) from existing registered devices this can be done using the [Manage registered devices](#) function. There is currently no fee to do this.

Follow the instructions in the [Updating Registrations](#) section of the **Device Registration Reference Guide** to [Manage registered devices](#).

## Update registered devices and products

If products need to be updated, for example Catalogue/Reference data has changed or was not added at the time of registration, this can be done using the [Update registered devices and products](#) function. If Medical Device Name or Model/Version needs to be updated, you will need to remove the product and add it again. There is currently no fee to do this.

If Basic UDI DI needs to be added follow the [Update Device Details](#) instructions in the **Device Registration Reference Guide**.

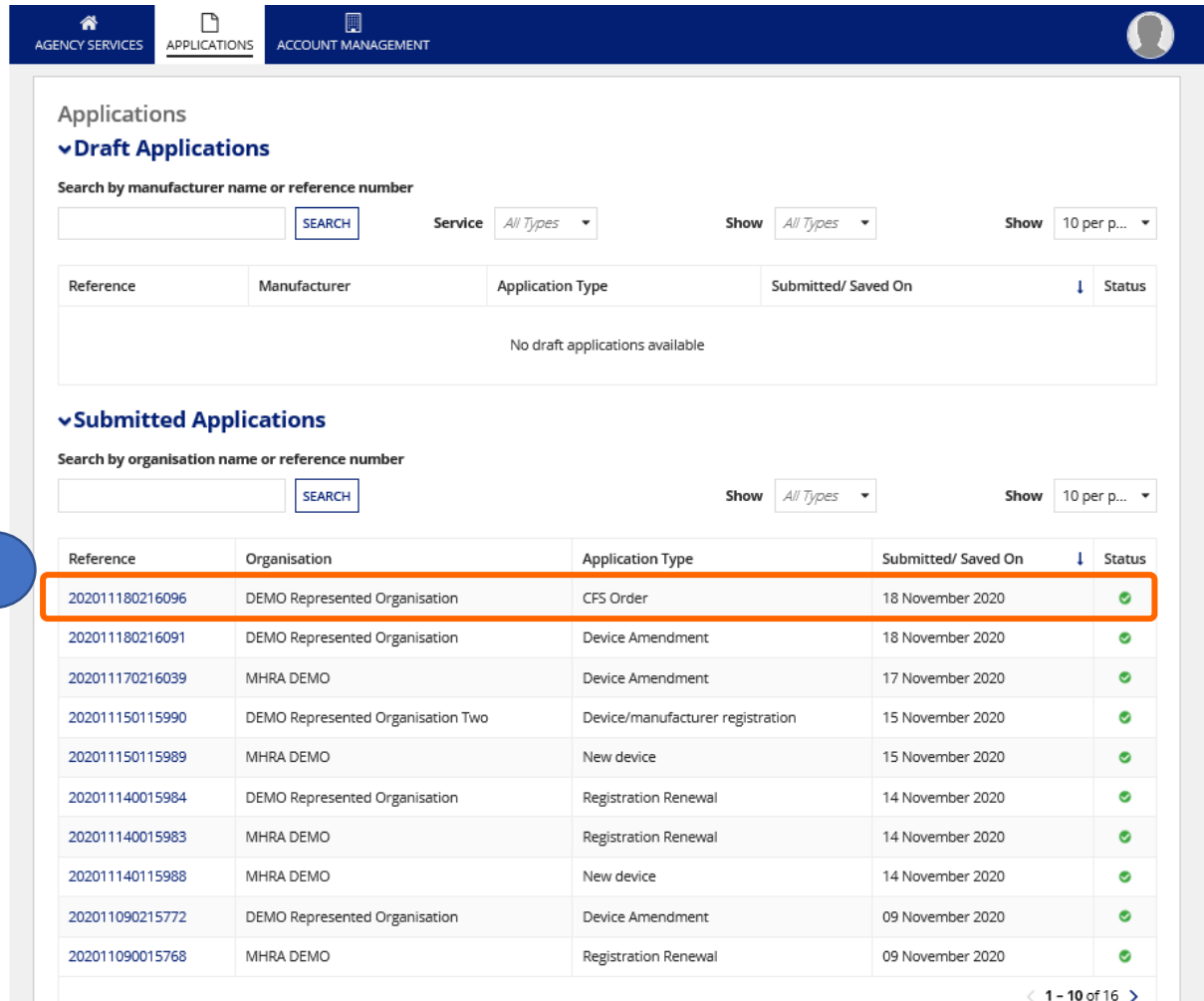
If product data needs to be updated, follow the [Update products](#) instructions in the **Device Registration Reference Guide**.

## Re-ordering a CFS

You can repeat a previous CFS order, optionally amending any of the information entered previously.

1. In your list of applications, **select** the previously completed application you would like to reorder by **clicking** on the **Reference number**.

**Please note** you will not be able to use the 'Reorder' functionality for any CFS orders submitted prior to 23 May 2018.

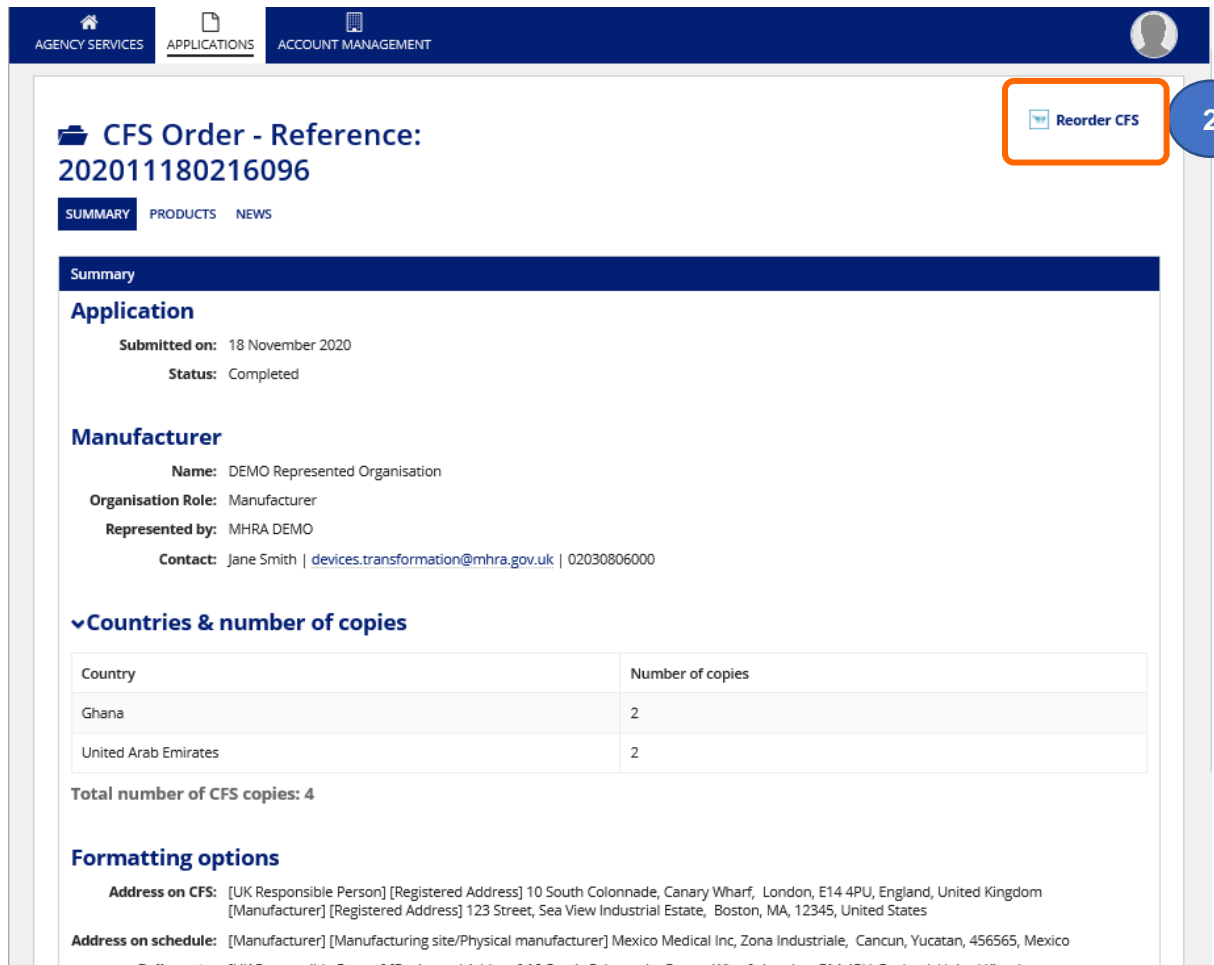


The screenshot shows the 'Applications' section of a web interface. It is divided into 'Draft Applications' and 'Submitted Applications'. The 'Submitted Applications' section contains a table with the following data:

Reference	Organisation	Application Type	Submitted/ Saved On	Status
202011180216096	DEMO Represented Organisation	CFS Order	18 November 2020	✓
202011180216091	DEMO Represented Organisation	Device Amendment	18 November 2020	✓
202011170216039	MHRA DEMO	Device Amendment	17 November 2020	✓
202011150115990	DEMO Represented Organisation Two	Device/manufacture registration	15 November 2020	✓
202011150115989	MHRA DEMO	New device	15 November 2020	✓
202011140015984	DEMO Represented Organisation	Registration Renewal	14 November 2020	✓
202011140015983	MHRA DEMO	Registration Renewal	14 November 2020	✓
202011140115988	MHRA DEMO	New device	14 November 2020	✓
202011090215772	DEMO Represented Organisation	Device Amendment	09 November 2020	✓
202011090015768	MHRA DEMO	Registration Renewal	09 November 2020	✓

At the bottom right of the table, there is a pagination indicator: < 1 - 10 of 16 >

2. Select the **Reorder CFS** link in the top right corner.



The screenshot shows the MHRA CFS Order Reference page for order 202011180216096. The page has a dark blue header with navigation tabs: AGENCY SERVICES, APPLICATIONS, and ACCOUNT MANAGEMENT. A user profile icon is in the top right. The main content area has a title 'CFS Order - Reference: 202011180216096' and a 'Reorder CFS' link highlighted with a blue circle and the number 2. Below the title are tabs for SUMMARY, PRODUCTS, and NEWS. The 'Summary' section is expanded, showing details for the Application, Manufacturer, and Countries & number of copies. The 'Countries & number of copies' section contains a table with 2 columns: Country and Number of copies. The 'Formatting options' section lists addresses for the CFS and the schedule.

**CFS Order - Reference:**  
202011180216096

**Summary** | PRODUCTS | NEWS

**Summary**

**Application**

**Submitted on:** 18 November 2020  
**Status:** Completed

**Manufacturer**

**Name:** DEMO Represented Organisation  
**Organisation Role:** Manufacturer  
**Represented by:** MHRA DEMO  
**Contact:** Jane Smith | [devices.transformation@mhra.gov.uk](mailto:devices.transformation@mhra.gov.uk) | 02030806000

**▼ Countries & number of copies**

Country	Number of copies
Ghana	2
United Arab Emirates	2

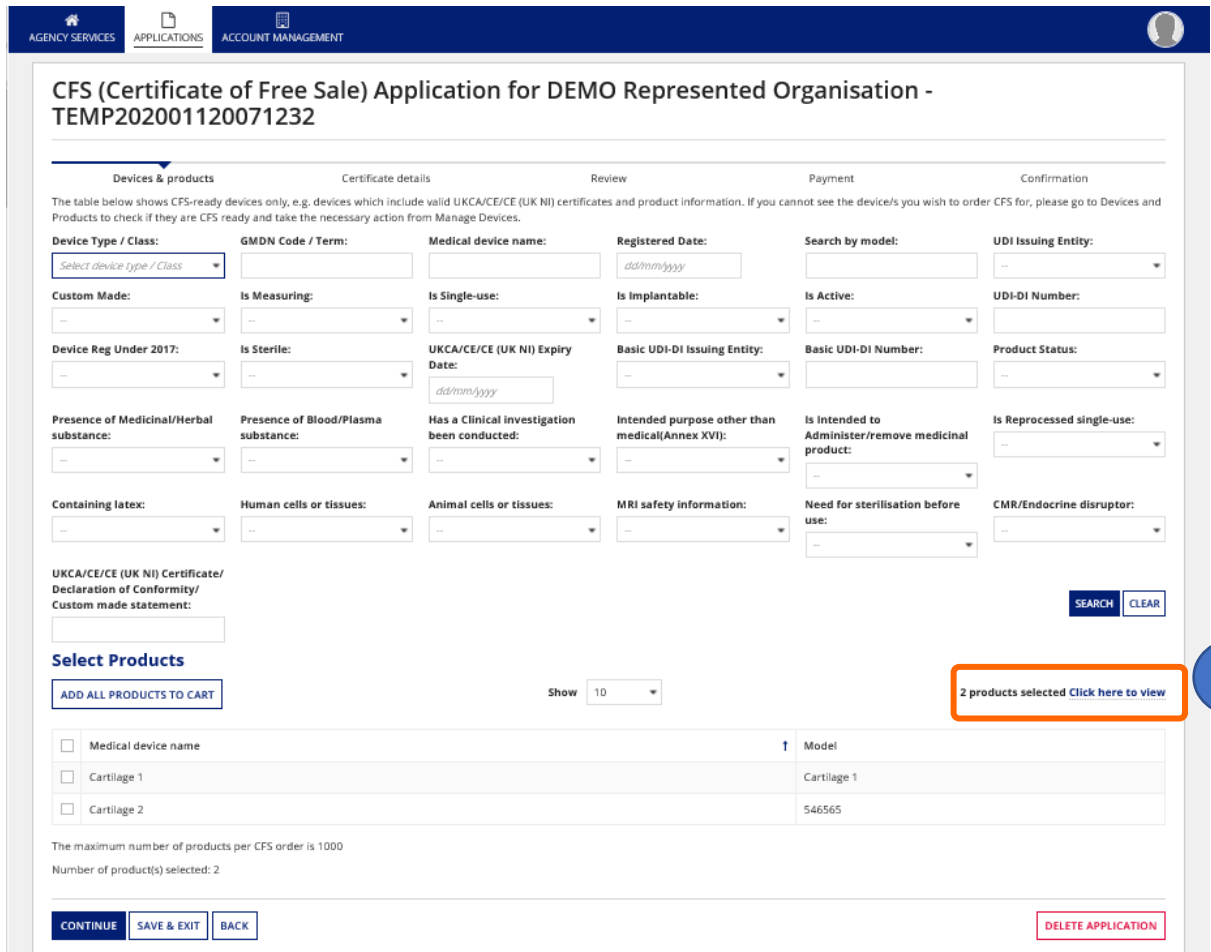
**Total number of CFS copies:** 4

**Formatting options**

**Address on CFS:** [UK Responsible Person] [Registered Address] 10 South Colonnade, Canary Wharf, London, E14 4PU, England, United Kingdom  
[Manufacturer] [Registered Address] 123 Street, Sea View Industrial Estate, Boston, MA, 12345, United States

**Address on schedule:** [Manufacturer] [Manufacturing site/Physical manufacturer] Mexico Medical Inc, Zona Industriale, Cancun, Yucatan, 456565, Mexico

- The products and information entered on the previous order will be pre-populated, providing Conformity Assessment certificates are still valid, but you can add or remove products for the new order as required. **Select** the [Click here to view](#) link to view all products on the original order.



**CFS (Certificate of Free Sale) Application for DEMO Represented Organisation - TEMP202001120071232**

The table below shows CFS-ready devices only, e.g. devices which include valid UKCA/CE/CE (UK NI) certificates and product information. If you cannot see the device/s you wish to order CFS for, please go to Devices and Products to check if they are CFS ready and take the necessary action from Manage Devices.

**Device Type / Class:**  **GMDN Code / Term:**  **Medical device name:**  **Registered Date:**  **Search by model:**  **UDI Issuing Entity:**

**Custom Made:**  **Is Measuring:**  **Is Single-use:**  **Is Implantable:**  **Is Active:**  **UDI-DI Number:**

**Device Reg Under 2017:**  **Is Sterile:**  **UKCA/CE/CE (UK NI) Expiry Date:**  **Basic UDI-DI Issuing Entity:**  **Basic UDI-DI Number:**  **Product Status:**

**Presence of Medicinal/Herbal substance:**  **Presence of Blood/Plasma substance:**  **Has a Clinical investigation been conducted:**  **Intended purpose other than medical(Annex XVI):**  **Is Intended to Administer/remove medicinal product:**  **Is Reprocessed single-use:**

**Containing latex:**  **Human cells or tissues:**  **Animal cells or tissues:**  **MRI safety information:**  **Need for sterilisation before use:**  **CMR/Endocrine disruptor:**

**UKCA/CE/CE (UK NI) Certificate/ Declaration of Conformity/ Custom made statement:**

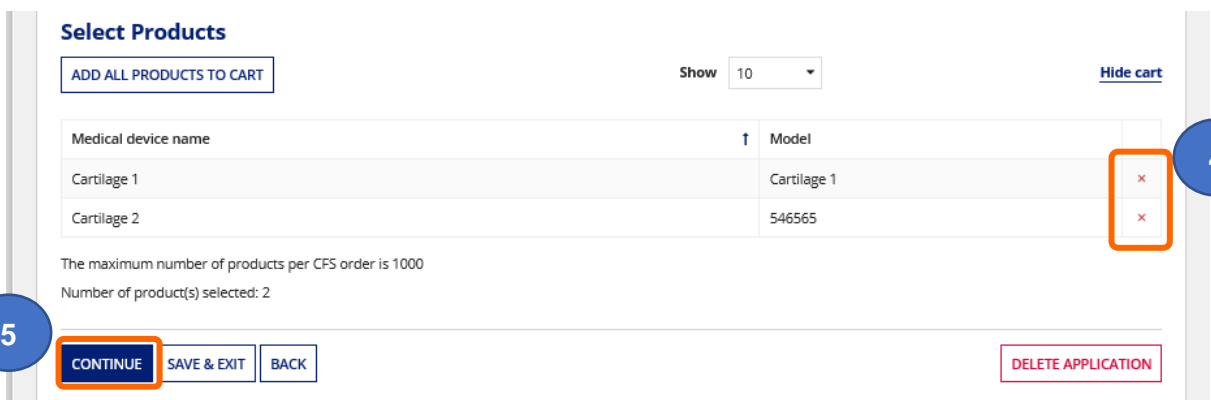
**Select Products**

**Show** 10

<input type="checkbox"/>	Medical device name	Model
<input type="checkbox"/>	Cartilage 1	Cartilage 1
<input type="checkbox"/>	Cartilage 2	546565

The maximum number of products per CFS order is 1000  
Number of product(s) selected: 2

- To remove product not required on this new order **click** the red **X** next to the product.



**Select Products**

**Show** 10

<input type="checkbox"/>	Medical device name	Model	
<input type="checkbox"/>	Cartilage 1	Cartilage 1	<input type="button" value="X"/>
<input type="checkbox"/>	Cartilage 2	546565	<input type="button" value="X"/>

The maximum number of products per CFS order is 1000  
Number of product(s) selected: 2

- Click the **Continue** button.

6. Make any required changes to [Addresses](#).

### CFS (Certificate of Free Sale) Application for DEMO Represented Organisation - TEMP202001120070701

---

Devices & products
Certificate details
Review
Payment
Confirmation

#### Address to be printed on CFS

**Registered Address:** 123 Street, Sea View Industrial Estate, Boston, MA, 12345, United States MANAGE ADDRESSES

**One additional address on certificates:**

[UK Responsible Person] [Registered Address] 10 South Colonnade, Canary Wharf, London, E14 4PU, England, United Kingdom

You can choose only one additional address to be printed on certificates.

**More addresses on schedule:**

[Manufacturer] [Manufacturing site/Physical manufacturer] Mexico Medical Inc, Zona Industriale, Cancun, Yucatan, 456565, Mexico

Any other addresses you choose will be printed on the schedule.

#### Special formatting request (optional)

Please note we will try to accommodate your request, but this cannot be guaranteed.

#### Certificate delivery address

- ● Registered address: 10 South Colonnade, Canary Wharf, London, E14 4PU, England, United Kingdom MANAGE ADDRESSES

#### Enter countries and number of certificates

Country	Number of certificates	
Ghana <span style="color: red; font-size: x-small;">✕</span>	2	<input type="checkbox"/> Do not specify a country <span style="color: red; font-size: x-small;">✕</span>

6

7

7. Make any required changes to [Countries and number of certificates](#). If you want to remove a country that was on the original order, **click** the red **X**.

8. Click continue to the [review](#) page and [Review your order and preview CFS Certificate and Schedule](#) then [Pay for your CFS Order](#).

**Important note:** Fee in screenshot is for illustrative purposes only. Check current [fee](#) on our website.

#### Enter countries and number of certificates

Country	Number of certificates	
Ghana <span style="color: red; font-size: x-small;">✕</span>	2	<input type="checkbox"/> Do not specify a country <span style="color: red; font-size: x-small;">✕</span>
United Arab Emirates <span style="color: red; font-size: x-small;">✕</span>	2	<input type="checkbox"/> Do not specify a country <span style="color: red; font-size: x-small;">✕</span>
<span style="color: #007bff; font-size: x-small;">➕ Add country</span>		

**Total number of certificates: 4**  
Price: £75.00

We charge £75.00 for 1 to 10 certificates and £10.00 for each additional certificate

CONTINUE

SAVE & EXIT

BACK

DELETE APPLICATION

8

## Save and exit: resume applications

When completing an application, you can save, exit and return to completing the application from where you left off. This option is not available on all screens.

Please follow the instructions in the **Device Registration Reference Guide**.

## Adding a New Manufacturer (for UKRP in UK and AR in NI)

Follow the instructions in the [Updating Registrations](#) section of **the Device Registration Reference Guide** for [Adding a Manufacturer](#). The current [statutory fee](#) will be payable.



## Annex I – Workflow

