

Attachment 4

Display of unapproved products under the Pharmaceutical Affairs Law

Similarly to ITEM2020, Web-ITEM2020 exhibitors are requested to comply with YAKUSEIHATSU No. 0609, June 9, 2017, No. 2, "Exhibition of unapproved medical devices at exhibitions, etc.". The following is a list of submissions, procedures and deadlines.

1. Items to be submitted

- (1) Unapproved medical device exhibition form (Attachment 4a) submitted as a PowerPoint file
- (2) Fill out the application form (Attachment 4b) with the required information in Word document.
- (3) Print the application form (Attachment 4b), affix the company seal, and scan the scanned PDF file.
- (4) Fill out the Medical Equipment Exhibit Request Form (Attachment 4c) as a Word document.

Please note that JIRA will combine (1) and the Exhibitor Request Form (4), which has the organizer's (JRC) seal on it. Please note that it will be posted on the website.

2. Procedures and deadlines

- (1) Send the above documents to JIRA by e-mail (deadline: April 24)
- (2) JIRA will compile the submitted documents and send them to the organizer (JRC), who will seal the exhibition request form, and then send a PDF of the request form to exhibitors by e-mail (around May 8).

Please note that even if the approval under the Pharmaceutical and Medical Device Act is granted by the end of the exhibition on June 5, the listings on Web-ITEM2020 cannot be changed.