Exhibitor Spotlight Sessions Rules and Regulations

February 12-16, 2021 – ORS 2021 Annual Meeting All Access

By applying for an Exhibitor Spotlight Session, a company agrees to adhere to all conditions and regulations outlined here. Whenever practical or appropriate, in the view of the ORS, disciplinary action will be progressive according to the violation of the listed Rules & Regulations. In the event of such restriction, the ORS will not be liable for any refunds on session costs or other exhibitor expenses.

Please be sure that your promotional department and anyone else involved in the arrangements of your session are aware of these Rules & Regulations. It is the responsibility of the Exhibitor to see that all involved in the creation of an Exhibitor Spotlight Session adhere to these rules.

CANCELLATIONS AND REFUNDS

Full payment is due at the time of application. Check payments must be received within 15 days of application in order to secure Spotlight Session. No refunds will be issued once payment is processed. The ORS reserves the right to reject or cancel any application for inability to follow rules and regulations, or for any reason by the ORS in its sole discretion.

FDA COMPLIANCE

All products that are not FDA approved for a particular use in humans or which are not commercially available in the United States will not be permitted to be shown in your session unless accompanied by the appropriate verbiage that indicate the device’s FDA clearance status. The following is the verbiage that should be displayed:

- This device is not cleared by the FDA for distribution in the United States.
- This device is intended to be used in the United States as described in the product’s labeling.

Verbiage must be easily visible and displayed when the device is being shown/demonstrated. The exhibitors shall have a letter from the FDA which describes the FDA clearance status of the designated use of the product or products. Exhibitors are cautioned about the FDA’s prohibition of promoting devices that are cleared for marketing for unapproved uses. Requests for information and guidance should be directed to:

FDA Division of Industry and Consumer Education (DICE) Center for Devices and Radiological Health (CDRH)
Phone: (800) 638-2041 / (301) 796-7100
Email: DI CE@fda.hhs.org

ORS LOGO

Use of the ORS logo is strictly prohibited. The Orthopaedic Research Society name, meeting name, and/or meeting logo may not be used without the explicit written permission from the ORS.

DEADLINES

Applications are due December 1, 2020. Sessions/videos must be submitted by January 4, 2021. Specific details on how and where to submit the session will be sent to you once you have confirmed participation.

REPRESENTATIVE FEES

Exhibitors will receive one (1) complimentary registration to ORS 2021 Annual Meeting All Access, with the purchase of a Spotlight Session.

ACCEPTANCE OF RULES AND REGULATIONS

By applying for an Exhibitor Spotlight Session, the exhibitor acknowledges receipt of and acceptance of these Rules & Regulations and all applicable rules of ORS 2021 Annual Meeting All Access as a condition of the ORS’s approval of Exhibitor Spotlight application. Discourteous language and other unacceptable behavior are a violation to these rules and will not be tolerated. The ORS may amend these Rules & Regulations at any time, and will inform exhibitors of such amendments in a timely manner.

ORTHOPAEDIC RESEARCH SOCIETY

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