EU MDR Compliance Declaration

REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017

Fisk Alloy does not manufacture any medical devices, and thus deems the requirements of Article 19 of EU MDR 2017/745, Annex IV not applicable to any of our products or services.

***Important Information**:* Information provided by Fisk Alloy on its Website or in other communications concerning the substance content of its products represents Fisk's knowledge and belief as of the date that it is provided. Fisk based its knowledge and belief on information provided by third parties and laboratory test results and makes no representation or warranty as to the accuracy of such information. Fisk has taken and continues to take reasonable steps to provide representative and accurate information but may not have conducted chemical analysis on all incoming materials. In no event shall Fisk Alloy, Inc. liability arising out of such information exceed the total purchase price of Fisk material sold by Fisk to a Customer on an annual basis.

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