

## BAUER Christophe

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**From:** Martina Giesemann [REDACTED]  
**Sent:** 18 February 2016 12:16  
**To:** EO-TriloguesConsultation  
**Subject:** [EOWEB] Trilogue on Medical Device Regulation

### Sender

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**Sender** Martina Giesemann [REDACTED]  
**To** Trilogues Consultation  
**Date** Thursday, February 18, 2016 12:15:42 PM CET

### Your data

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#### Part 1 - Contact information

<b>First name</b>	Martina
<b>Surname</b>	Giesemann
<b>Gender</b>	Female
<b>E-mail address</b>	[REDACTED]
<b>Language you would like to receive an answer in</b>	en - English
<b>Other language you would accept an answer in (if applicable)</b>	de - Deutsch

#### Part 2 - Data

**To** Trilogues Consultation  
**Subject** Trilogue on Medical Device Regulation  
Dear Madam,

My company is acting as EU Auth. Rep. for medical device manufacturers. As such, it is extremely important for us to keep up to date with regulatory changes. Since the trilogue started last year, we found it very difficult to do this. Information were very rare. This makes it near impossible for us to prepare our clients (the medical device manufacturers) of the forthcoming changes.

**Content** Thank you.

Regards

Martina Sander-Giesemann

mdi Europa GmbH