

TO WHOM IT MAY CONCERN

Your reference	Our reference	Tel. +49/56 61/71-	Fax +49/56 61/71-	Date
	HC-RA-DE08E JE/KJ	18 70	18 70	June 14, 2012

CONFIRMATION

This is to confirm that our medical devices

<i>Product</i>	<i>Art. no.</i>
Dosifix®	4037014
Exadrop®	4061209
Exadrop®	4061225
Exadrop®	4061284
Exadrop®	4061306
Exadrop®	4062264
Exadrop®	4180330
Intrafix® Primeline	4060563
Intrafix® Primeline	4062191
Intrafix® Primeline	4062957
Intrafix® Primeline	4063287
Intrafix® Primeline	4062981L
Intrafix® SafeSet	4063000
Intrafix® SafeSet	4063001
Intrafix® SafeSet	4063002
Intrafix® SafeSet	4063004
Medifix®	4276108
Medifix®	4276116
Medifix®	4276612
Medifix®	4276620
Medifix®	4276728
Sangofix®	4107411

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<i>Product</i>	<i>Art. no.</i>
Sangofix®	4117301
Sangofix®	4117417F

are DEHP-free (free of Di(2-ethylhexyl)phthalat).

For and on behalf of

B. Braun Melsungen AG

i. V.


Ulrich Jedelhauser
Manager Regulatory Affairs CoE IV-Systems / IV-Access

i. A.


Kirsten Janssen
Administrator Regulatory Affairs CoE IV-Systems