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## Data in Brief

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## Data Article

Data on the fate of MACS<sup>®</sup> MicroBeads intramyocardially co-injected with stem cell productsPaula Müller<sup>a,b,1</sup>, Ralf Gaebel<sup>a,b,1</sup>, Heiko Lemcke<sup>a,b</sup>,  
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## Keywords:

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Haematopoietic stem cells (HSCs)

Mesenchymal stem cells (MSCs)

Magnetic activated cell sorting (MACS<sup>®</sup>)MACS<sup>®</sup> MicroBeads

## ABSTRACT

The data presented in this article are related to the research article “Intramyocardial Fate and Effect of Iron Nanoparticles co-injected with MACS<sup>®</sup> purified Stem Cell Products” (Müller et al., 2017) [1]. This article complements the cellular localization of superparamagnetic iron dextran particles (MACS<sup>®</sup> MicroBeads) used for magnetic activated cell sorting (MACS<sup>®</sup>). Data evaluate the time-dependent detachment of these nanoparticles from CD133<sup>+</sup> haematopoietic stem cells (HSCs) and CD271<sup>+</sup> mesenchymal stem cells (MSCs). Furthermore, the influence of these stem cells as well as of nanoparticles on cardiac remodeling processes after myocardial infarction (MI) was investigated.

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## Specifications Table

Subject area	Biology
More specific subject area	Intramyocardial transplantation of MACS <sup>®</sup> purified stem cell products
Type of data	Image, graph, figure, text file
How data was acquired	Structured illumination microscopy (Zeiss ELYRA PS.1 LSM 780), flow cytometry (BD LSR-II), histological staining
Data format	Analyzed
Experimental factors	CD133 <sup>+</sup> and CD271 <sup>+</sup> stem cells were automatically (using the CliniMACS <sup>®</sup> Prodigy BM-133 system) and manually (using Mini MACS <sup>®</sup> technology) isolated from human bone marrow (BM)
Experimental features	Investigation of the Intracellular localization and time-dependent detachment of MACS <sup>®</sup> MicroBeads from stem cells using the Labeling Check Reagent-FITC (Miltenyi Biotec). Impact of MACS <sup>®</sup> MicroBeads on collagen deposition after myocardial infarction using an ischemia/reperfusion mouse model and Sirius Red staining.
Data source location	Rostock University Medical Center, Schillingallee 69, 18057 Rostock, Germany
Data accessibility	The data are available with this article

## Value of the data

- MACS<sup>®</sup> is the most commonly used technique for the purification of stem cell subpopulations intended for the treatment of cardiovascular diseases.
- Data about the binding of MACS<sup>®</sup> MicroBeads to stem cells are crucial for *in vivo* application of stem cell products.
- Data provide information about the effect of co-injected MACS<sup>®</sup> MicroBeads on cardiac remodeling processes after MI.
- Data can be useful for other researchers analyzing the cardiac regeneration potential of MACS<sup>®</sup> purified stem cells products.
- Data clarifies the safety of MACS<sup>®</sup> MicroBeads for clinical application.

## 1. Data

The data include information about the cellular localization of MACS<sup>®</sup> MicroBeads (labelled with Labeling Check Reagent-FITC) right after the manual MACS<sup>®</sup> based isolation of CD133<sup>+</sup> and CD271<sup>+</sup> stem cells (Fig. 1). The detachment of FITC-labelled MACS<sup>®</sup> MicroBeads was evaluated by measuring the time-dependent fluorescence intensity of MACS<sup>®</sup> purified CD133<sup>+</sup> cells incubated under cell culture conditions (37 °C in StemSpan<sup>™</sup> H3000) using flow cytometry (Fig. 2). Furthermore, the effect of manually and automatically (Good Manufacturing Practice (GMP)-conform) MACS<sup>®</sup> purified CD133<sup>+</sup> and CD271<sup>+</sup> stem cells as well as of MACS<sup>®</sup> MicroBeads on fibrosis after MI was assessed in a cardiac ischemia/reperfusion mouse model by histological staining (Fig. 3).

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