

Declaration of Biological Shipments for RESEARCH

You need: Selection 2:	 an invoice stating: The correct name and origin (source) of the reagents The number of vials / or other items the packaging must be labeled
Selection 2:	and packaging made be labeled
	highly purified antibodies.
	Fill in the correct name (i.e., goat anti mouse XYZ) $\ \square$ monoclonal $\ \square$ polyclonal
state:	the antibody was separated from plasma/serum or ascites fluid and highly purified so that animal disease or other pathogenic agents were killed off or effectively removed.
if not	see Selection 4 and 5
Selection 3:	Cell cultures, Cell lines, Fill in the correct name:
in:	\square serum free medium \square commercial medium with serum (see Selection 4 and 5)
Selection 4:	\square Blood / Serum \square Tissue \square Organs \square Other
a.) fixated in:	☐ Alcohol
b.) untreated:	the <u>consignee</u> will have had to apply for a special derogation
state:	The materials sent
	are non- infectious research material, do not contain any pathogenic agents.
	do not contain any pathogenic agentswere never contaminated by any infectious material.
Selection 5:	The material will be sent in ready-made, commercial serum
	□ yes: Cat No produced by
	Product Data-Sheet has to be attached
	\square no: the <u>consignee</u> will have had to apply for a special derogation
Selection 6:	otherwise not named biological specimens: □ pathogenic □ apathogenic
	Please provide scientific name:
	Intended Use:
If you do n	ot have a stamp, put your statements on a separate letterhead
ease attach this f	form on outer packaging. This information is correct and true.



In-Vitro laboratory reagents COMMERCIAL use

Please supply the necessary documentation to ensure timely veterinary clearance of animal derived laboratory reagents to Germany.

Selection 1: any Biological Product of <u>ANIMAL</u> origin.

You need: an invoice stating: Consignee and Consignor

- The correct name and origin (source) of the reagents given in the invoice
- The number of receptacles / net weight in milligram are required for all products
- physical products must be labeled
- and marked for their purposes i.e.: "for in-vitro research"

For In-vitro Diagnostics / Medical Devices (CE approved):

You need: a statement:

all the items contained in the shipment:

AirwayBillNumber:

meet the criteria of in-vitro diagnostics /medical devices following Directive 98/79/EC or Directive 93/42/EWG

manufacturer: i.e Cell Marque authorized representative: i.e. Emergo group

For intermediate products, products intended for further manufacturing:

You need: The proper **Declaration** following the Regulation EC No. 142/2011

• This certificate will be issued by the consignee.

- Please make sure <u>you are listed</u> as approved for the import into the EC.
- Please make sure the consignee is listed as approved

For untreated blood products:(i.e. Serum, BSA)

You need: The proper **health certificate** following the Regulation

EC No. 142/2011

For treated blood products: (i.e. Serum, BSA)

You need: The proper **health certificate** following the Regulation

EC No. 142/2011

For ready made laboratory reagents:

You need: Provided they are none of the above, an **invoice** (see selection 1)

But they have to be ready made for the EU common market,

including the proper labelling