

BRIEF ORIGINAL

# English

# Long-term stability of trastuzumab in plasma and whole blood samples stored under different conditions

Estabilidad a largo plazo del trastuzumab en plasma y suero almacenados bajo diferentes condiciones

Jonathan González-García<sup>1</sup>, Fernando Gutiérrez-Nicolás<sup>1</sup>, Gloria Julia Nazco-Casariego<sup>1</sup>, María Micaela Viña-Romero<sup>2</sup>

<sup>1</sup>Servicio de Farmacia, Complejo Hospitalario Universitario de Canarias, San Cristóbal de La Laguna, Tenerife. Spain. <sup>2</sup>Servicio de Farmacia, Hospital Universitario Nuestra Señora de La Candelaria, Santa Cruz de Tenerife, Tenerife. Spain.

#### Author of correspondence

Fernando Gutiérrez Nicolás Complejo Hospitalario Universitario de Canarias. Ctra. General La Cuesta s/n (38320), San Cristóbal de La Laguna, Tenerife. Spain.

Farmacia Hospitalaria | ONLINE FIRST ||

Email: fgunico@hotmail.com

: DOI: 10.7399/fh.11110

# Abstract

**Objective:** The present work analyzed the stability of trastuzumab concentrations in blood and plasma samples stored at 4 °C and -20 °C.

**Method:** Blood samples of patients treated with trastuzumab (Herceptin®) were analyzed. Stability of trastuzumab was analyzed under different conditions: whole blood stored at 4 °C during 72 hours, blood plasma stored at 4 °C during 120 hours and blood plasma stored at -20 °C during 365 days.

**Results:** In whole blood trastuzumab concentration at 72 hours was 99.01  $\pm$  0.02%, and in plasma samples at 120 hours it was 98.6  $\pm$  2.0%. The concentration of trastuzumab was 95.22  $\pm$  3.20% after 12 months' storage at -20 °C.

**Conclusions:** The concentration of trastuzumab remains stable for at least 72 hours in whole blood stored at 4 °C, five days in plasma stored at 4 °C and one year in plasma stored at –20 °C.

## Resumen

**Objetivo:** Analizar la estabilidad del trastuzumab en sangre y plasma a 4 °C y -20 °C.

**Método:** Se determinó la concentración plasmática de trastuzumab en las muestras de sangre y plasma de pacientes con cáncer de mama HER2 positivo tratados con trastuzumab (Herceptin®). Se analizó la estabilidad del trastuzumab en sangre a 4 °C durante 72 horas. En el caso de las muestras de plasma, se estudió la estabilidad a 4 °C durante 72 y 120 horas, así como durante 365 días congeladas a -20 °C.

**Resultados:** En sangre, la concentración de trastuzumab a las 72 horas fue de 99,01  $\pm$  0,02%, y en las muestras de plasma a las 120 horas fue de 98,6  $\pm$  2,0%. La concentración de trastuzumab fue de 95,22  $\pm$  3,20% después de 12 meses de almacenamiento a -20 °C.

**Conclusiones:** El trastuzumab permanece estable durante al menos 72 horas en sangre total almacenada a 4 °C, cinco días en plasma almacenado a 4 °C y un año en plasma almacenado a –20 °C.

#### **KEYWORDS**

Blood plasma; Blood serum; Stability; Storage; Trastuzumab.



**PALABRAS CLAVE** Almacenamiento; Estabilidad; Plasma; Suero; Trastuzumab. Los artículos publicados en esta revista se distribuyen con la licencia Articles published in this journal are licensed with a Creative Commons Artichulion-NonCommercial-ShareAlike 4.0 International License. http://creativecommons.org/licenses/by-nc-sa/4.0/ La revista Farmacia no cobra tasas por el envis de trabajos, ni tampoco por la publicación de sus artículos.

#### Introduction

Trastuzumab is a monoclonal antibody used in the treatment of breast and gastric cancer where neoplastic tissue overexpresses the epidermal growth factor receptor HER2<sup>1,4</sup>. Recently, a link has been established between trough levels (C<sub>min</sub>) of trastuzumab and effectiveness<sup>5,8</sup>, which has attracted interest in monitoring the pharmacokinetics of this drug.

One of the prerequisites for pharmacokinetic study is to determine the stability of the drug in blood samples, because sometimes they cannot be processed immediately after extraction and are stored. To date, there are no data on the long-term stability of trastuzumab in whole blood and/or plasma stored under different conditions. The present work analyzes the stability of trastuzumab concentrations in blood and plasma samples stored at 4 °C and -20 °C, during differents points of time (24, 72, 120 hours and one year).

### Methods

#### Study design and sample collection

We performed a single-center, prospective, observational study of a year of duration, in which blood samples (5 mL) of HER2 positive breast cancer patients treated with trastuzumab (Herceptin®) were analyzed. The samples were collected in K<sub>2</sub>EDTA tubes (Becton Dickson Company®, USA) and centrifuged at 3,000 g for 5 minutes to obtain blood plasma.

We analyzed the stability of the samples stored under different conditions:

- Analysis 1: whole blood stored at 4 °C during time zero (CO), at 24 and 72 hours.
- Analysis 2: blood plasma stored at 4 °C during C0, 24, 72 and 120 hours.
- Analysis 3: blood plasma stored at -20 °C, during C0, 60, 180 and 365 days.

Trastuzumab concentration measured at time zero (CO), i.e. immediately after extraction, was considered the reference value; all determinations were

performed in duplicate. The results are expressed as mean  $\pm$  standard deviation (SD) for both.

Statistical analysis was performed using IBM SPSS v22 (IBM Corporation, Armonk, NY).

#### Compliance with ethical standards

The present analysis was part of a wider study (EPA-SP with code FER-TRA-2015-01), approved by the Clinical Research Ethics Committee of our hospital for implementation in February 2015. Written informed consent was obtained from all patients before inclusion in the study.

#### Determination of trastuzumab concentrations

Plasma concentrations of trastuzumab were determined using ELISA (enzyme-linked immunosorbent assay), following the manufacturer's specifications (Shikari® q-tras)°, employing ELISA triturus autoanalyzer (Grifols®). The samples were diluted 20 or 80 times (depending on the estimated concentration) with buffer solution (commercial kit) in duplicate to minimize pipetting errors and intrinsic variations of the method.

#### Stability of trastuzumab

For the analysis of the stability of trastuzumab in blood samples, the drug concentration calculated at time zero (CO) was taken as the reference value. Thus the stability of the drug in samples stored for different times was expressed as a percentage of the CO value. A stability limit of 90% was established as acceptable.

For the analysis of stability at 4 °C (whole blood and plasma) we used three samples (A, B and C) and for the study of stability at –20 °C (plasma) we used five samples (D, E, F, G and H).

#### Results

A total of 82 determinations were made, with the following results:

Table 1. Mean concentrations of trastuzumab in whole blood and plasma samples stored at 4 °C for periods ranging from 0 to 120 hours

	Mean concentration µg/mL (± SD)								
	Sam	ple A	Sam	ple B	Sample C				
	Blood	Plasma	Blood	Plasma	Blood	Plasma			
C0	34.6 (0.9)	34.7 (0.3)	24.8 (1.8)	25.4 (0.3)	46.3 (.0)	46.1 (0.4)			
24 hours	35.5 (1.8)	33.3 (0.6)	26.0 (0.2)	26.2 (0.3)	46.5 (0.5)	47.7 (0.6)			
72 hours	36.1 (0.4)	33.9 (0.6)	24.4 (0.1)	26.7 (0.6)	47.1 (0.2)	44.6 (0.5)			
120 hours	ND	33.3 (0.1)	ND	26.1 (0.3)	ND	44.5 (0.3)			
% <b>CO</b>	100	95.7	98.1	97.2	98.4	96.6			

CO: Concentration of trastuzumab measured immediately after blood sample extraction (time zero); ND: not data.

Table 2. Mean concentrations of trastuzumat	b in plasma samples stored at –2	20 °C for periods ranging from 0 to 12 months
---	----------------------------------	---

	Sample D		Sample E		Sample F		Sample G		Sample H	
	Conc. (± SD)	% C0								
CO	46.2 (0.1)	-	27.2 (1.9)	-	37.7 (0.6)	-	111.0 (1.1)	-	62.6 (0.3)	-
2 months	48.4 (0.1)	95.2	26.7 (0.8)	98.3	38.2 (0.1)	98.5	103.9 (6.2)	93.6	62.1 (1.1)	98.9
6 months	46.5 (1.1)	99.2	28.4 (0.9)	95.4	36.4 (0.3)	96.5	111.3 (1.4)	99.8	59.9 (0.6)	95.3
12 months	45.3 (1.5)	98.2	24.9 (0.3)	91.5	34.6 (0.6)	91.9	108.1 (1.1)	97.3	61.1 (0.4)	97.2

CO: plasma concentration of trastuzumab measured immediately after blood sample extraction (time zero). Conc.: concentration (µg/ml).

#### Stability of trastuzumab in samples stored at 4 °C

The mean concentrations of trastuzumab are shown in Table 1. In whole blood at 72 hours this was 99.01  $\pm$  0.02, and in plasma samples at 120 hours it was 98.6  $\pm$  2.0% (Table 1).

# Stability of trastuzumab in plasma samples stored at -20 °C

The mean concentration of trastuzumab was 96.9  $\pm$  1.7%; 97.2  $\pm$  0.9% and 95.22  $\pm$  3.20%, after two, six and 12 months' storage at –20 °C respectively (Table 2).

#### Discussion

Monitoring plasma levels of drugs has been shown to be an effective tool to improve clinical outcomes of treatments<sup>10</sup>. However, this is not always easy, as there are often limitations such as the processing and storage of samples<sup>11</sup>. Jamieson *et al.*<sup>o</sup> have shown that trastuzumab is stable after 72 hours and 10 weeks in plasma samples stored at 4 °C and –20 °C, respectively. With this work we wished to show that sample storage time can exceed 120 hours at 4 °C and that plasma samples can even be stored for one year at –20 °C. The present study not only analyzed trastuzumab

## Bibliography

- Bang YJ, Van Cutsem E, Feyereislova A, Chung HC, Shen L, Sawaki A, et al. Trastuzumab in combination with chemotherapy versus chemotherapy alone for treatment of HER2-positive advanced gastric or gastro-oesophageal junction cancer (ToGA): a phase 3, open-label, randomised controlled trial. Lancet. 2010;376(9742):687-97.
- Slamon DJ, Leyland-Jones B, Shak S, Fuchs H, Paton V, Bajamonde A, et al. Use of chemotherapy plus a monoclonal antibody against HER2 for metastatic breast cancer that overexpressed HER2. N Engl J Med. 2001;344(11):783-92.
- Cobleigh MA, Vogel CL, Tripathy D, Robert NJ, Scholl S, Fehrenbacher L, et al. Multinational study of the efficacy and safety of humanized anti-HER2 monoclonal antibody in women who have HER2-overexpressing metastatic breast cancer that has progressed after chemotherapy for metastatic disease. J Clin Oncol. 1999;17(9):2639-48.
- Vogel CL, Cobleigh MA, Tripathy D, Gutheil JC, Harris LN, Fehrenbacher L, et al. Efficacy and safety of trastuzumab as a single agent in first-line treatment of HER2overexpressing metastatic breast cancer. J Clin Oncol. 2002;20(3):719-26.
- Cosson VF, Ng VW, Lehle M, Lum BL. Population pharmacokinetics and exposureresponse analyses of trastuzumab in patients with advanced gastric or gastroesophageal junction cancer. Cancer Chemother Pharmacol. 2014;73(4):737-47.

stability in blood plasma, but also confirmed its stability in whole blood samples stored for 72 hours at 4 °C. These data are of particular interest to health centers that lack the technical means for processing the samples which are then sent to external laboratories.

In conclusion, the concentration of trastuzumab remains stable for at least 72 hours in whole blood stored at 4 °C, five days in plasma stored at 4 °C and one year in plasma stored at –20 °C.

#### Funding

No funding.

#### **Conflict of interests**

No conflict of interests.

## Contribution to the scientific literature

The stability of trastuzumab in blood or plasma has not been assessed. Such assessment is fundamental to conducting pharmacokinetic studies. The present study shows that the drug is stable in serum for 5 days when stored between 0°C and 5°C and up to 1 year when stored at -20°C.

- Kyi C, Shah MA. A case report of trastuzumab dose in gastric cancer. J Gastrointest Oncol. 2013;4(4):E19-22.
- 7. González-García J, Alonso-Álvarez B, Nazco-Casariego GJ, Batista-López N, Gutiérrez-Nicolás F. Plasma levels of trastuzumab in gastric cancer: Case report. J Oncol Pharm Pract. 2017;23(8):635-7.
- González García J, Gutiérrez Nicolás F, Nazco Casariego GJ, Batista López JN, Ceballos Lenza I, Ramos Díaz R, et. al. Influence of Anthropometric Characteristics in Patients With Her2-Positive Breast Cancer on Initial Plasma Concentrations of Trastuzumab. Ann Pharmacother. 2017;51(11):976-80.
- Jamieson D, Cresti N, Verrill MW, Boddy AV. Development and validation of cellbased ELISA for the quantification of trastuzumab in human plasma. J Immunol Methods. 2009;345(1-2):106-11.
- Lesko U, Schmidt S. Individualization of drug therapy: history, present state, and opportunities for the future. Clin Pharmacol Ther. 2012;92(4):458-66.
- Ramos-Díaz R, Gutiérrez-Nicolás F, Nazco-Casariego GJ, González-Perera I, Pérez-Pérez JA. Validation of a fast and low-cost alkaline lysis method for gDNA extraction in a pharmacogenetic context. Cancer Chemother Pharmacol. 2015;75(5):1095-8.