

Determination of Ganciclovir in Plasma of Newborns with Congenital CMV Infection

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ABSTRACT

Infections caused by human cytomegalovirus are important causes of fetal and neonatal morbidity and mortality. Ganciclovir ([[(9-(1,3-dihydroxy-2-propoxymethyl) guanine, GCV) is a synthetic acyclic nucleoside which has shown activity against Cytomegalovirus. GCV treatment has been associated with serious toxic hematological effects such as neutropenia and leukopenia, thus drug monitoring is needed, especially in the case of newborns. The aim of the work is to develop and validate an HPLC method for the quantification of GCV in plasma.

Abbreviations: GCV: Ganciclovir; VGCV: Valganciclovir; HPLC: High-Performance Liquid Chromatography; RT: Retention Time; AUC: Area Under The Concentration Time Curve; LLOQ: Lower Limit Of Quantifications; LLOD: Lower Limit Of Detection; SD: Standard Deviation; RE: Relative Error; CV: Coefficient Of Variation

Introduction

The transmission of cytomegalovirus (CMV) to the fetus is a result of maternal viremia and transplacental infection often maternal CMV infections are subclinical. Mortality in symptomatic infants is about 10%. Surviving newborns can contract mental retardation and other neurologic deficits [1,2]. The treatment of congenitally cytomegalovirus infected neonates involves the use of ganciclovir or its prodrug valganciclovir (GCV). Unfortunately, their use is limited by neutropenia, carcinogenicity, gonadal dysgenesis risks, development of viral resistance and drug-drug interactions [3,4]. Several strategies have been suggested to prevent high plasma ganciclovir concentrations and toxicity. Dosage reduction is recommended in patients with renal damage. Temporary discontinuation of an interacting medication, adjustment of dosages and careful monitoring of the patient is suggested to prevent any potential adverse effects of such interactions [5].

Material and Method

Chemicals

Ganciclovir and Trifluoroacetic acid (TFA) were purchased from Sigma Chemical Co. (St. Louis, MO). and 1-heptanesulfonic

acid. Distilled water and perchloric acid were purchased from Sigma-Aldrich Chemical (St. Louis, MO, USA).

Blood Samples Preparation

Blood samples (1.5 ml) were collected in gel separating tubes. The blood was then centrifuged for 5 min at 13000 rpm. 1 ml of the serum thus obtained were added with 250 μ l of TFA 20 % and the suspension was then centrifuged at 13000 rpm for 10 min. Aliquots of deproteinated serum were stored at -80°C until use.

GCV Extraction

100 μ l of serum was added with 95 μ l of distilled water and 50 μ l of perchloric acid 7 %.

Assay Application

8 newborns with congenital CMV infection were treated with 15 mg/kg of Valganciclovir (GCV prodrug) twice daily. Blood samples were collected before (T0) and after 2 hours (T1) of the morning administration.

Chromatographic Conditions

The chromatographic system consisted of an HPLC Jasco LC-Net II/ADC (JASCO International Co., Ltd. Tokyo, Japan) with a 20

μ L Rheodyne 8125 injector (Rheodyne, Ronherth Park, CA, USA), an UV Detector UV "Jasco MD-2010 plus" (JASCO International Co., Ltd. Tokyo, Japan) (σ 254nm), and a reversed phase Jupiter C-18 5 σ m (150x4.6 mm) column (Phenomenex, California, USA). Using Distilled water as mobile phase and 1.2 ml/min flow rate.

Results

Recovery of GCV

Recovery of ganciclovir from the extraction process was calculated by using the next formula:

$$\text{Recovery\%} = \frac{C_1}{C_2} \times 100$$

C^1 = analyte concentration measured after addition

C^2 = concentration of added standard

Results are reported in Table 1.

Table 1: Recovery.

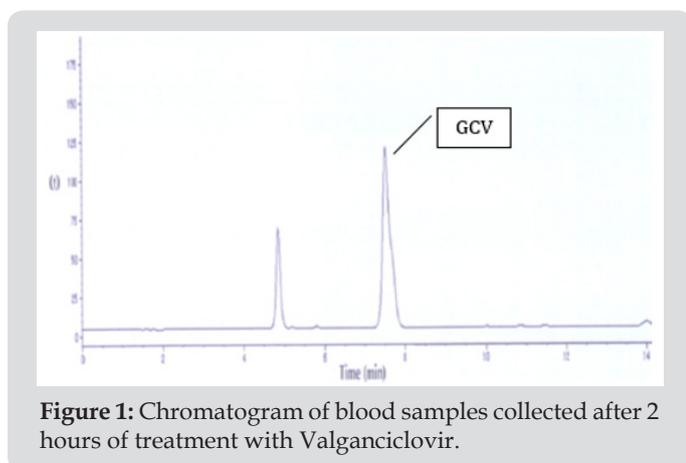
The Amount Added (mg)	Recovery (%)	CV (%)
0.23	110.51	1.87
0.29	102.82	3.56
0.33	104.97	2.39

GCV Stability

GCV concentrations were stable in standard solutions after one month of storage at 4°C, in plasma for at least 5 h at room temperature, in biological matrices when stored at 220°C for at least six months.

Method Validation

The method was validated, considering linearity, LLOD, LLOQ, precision and accuracy, according to European Medicines Agency guidelines Validation [6]. The method of constant addition was used for the construction of calibration curves Figure 1.

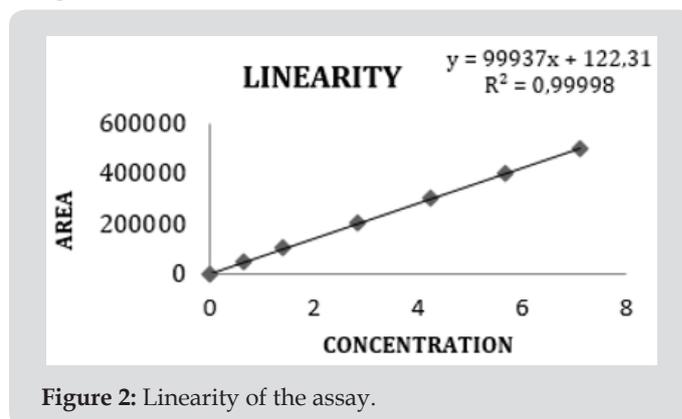


Method Selectivity

The selectivity of the assay was defined by analysis of blank plasma, with and without internal standard. There are no interfering peaks present at the retention time for GCV peaks.

Method Linearity

The correlation coefficient of the linear regression was reported in Figure 2.



Method Sensitivity

The method detection limit was tested by repeated analysis of blank samples.

The detection limits were determined as the concentration giving a peak height three times the noise background

Lower limit of Detection (LLOD) = $3,3 \sigma / S = 0,021 \mu\text{g} / \text{mL}$

Lower Limit of Quantitation (LLOQ) = $10 \sigma / S = 0,065 \mu\text{g} / \text{mL}$

σ = DS of the y intercept of regression line

S=slope of regression line

The quantitation limit of an individual analytical procedure is the lowest amount of analyte in a sample which can be quantitatively determined with suitable precision and accuracy. LLOQ (0.065 $\mu\text{g} / \text{ml}$), was >10

Method Accuracy

The accuracy of the method was tested by relative error calculation. The accuracy values in intra-day variation studies at low, medium and high concentrations of GCV in plasma were within acceptable limits (n=5) Table 2.

Table 2: Relative error & Relative standard deviation of regression line slope.

Nominal Value(μ g/ML)	Fitted Value(μ g/ML)	RE%	RSD%
0.1	0.10 \pm 0.01	8.57	11.4
0.5	0.51 \pm 0.02	3.39	3.35
1	1.00 \pm 0.02	1.15	1.83
2	1.98 \pm 0.02	0.96	0.87
5	5.01 \pm 0.05	0.69	0.92

Acceptable limits:<15%

Method Precision

The precision of the method was expressed by the coefficient of variation of intra- and inter-day variations of the assay under

the same operating conditions. The acceptance criterion was set at 15% Table 3.

Table 3: Intra and Inter-day variability and stability.

Conditions	24 Hours	7 Days
Room temperature	1.17	3.05
4°C	3.28	6.39
-20°C	0.22	18.36

Method Applicability

Applicability of the method was confirmed by analyzing samples from 8 newborns with congenital CMV infection treated with 15 mg/kg of Valganciclovir (GCV prodrug) twice daily. GCV concentration in blood samples collected before (T₀) and after 2 hours (T₁) of the morning administration are reported in Table 4.

Table 4: Assay application.

Subject	T ₀ (Mg/MI)	T ₁ (Mg/MI)
A	0.21±0.09	0.87±0.06
B	0.42±0.00	1.04±0.10
C	0.28±0.02	1.98±0.19
D	0.21±0.13	2.04±0.13
E	0.11±0.02	1.99±0.02
F	0.06±0.06	2.00±0.77
G	0.35±0.37	2.70±0.03
H	0.01±0.01	1.94±0.10

The present HPLC method for the determination of GCV was useful for the therapeutic monitoring of antiviral medication.

Discussion

Amperometry detection [7], fluorescence detection, ion-pairing agents [8,9], and UV detection [10-13], are general used to determine GCV. The difficult to detect GCV in newborn plasma are small volume of plasma available and little GCV concentration. The new analytical method developed was validated according to European Medicines Agency guidelines Validation. For the validation of the extraction procedures, samples of blank serum were spiked with known amounts of GCV. The recovery was determined by computing the ratio of the amount extracted from spiked samples to the amount added, determined by the injection of spiked solution. Ganciclovir was completely recovered during the extraction. Good ability to obtain results directly proportional to the concentration of analyte (not more than some number close to 1), fine detection limit, high precision demonstrated by very good intra- and inter-day repeatability (mean recoveries were near 100% and %RSD values were low) make this method a useful tool for the therapeutic monitoring of GCV medication. Applicability of the method was confirmed by analyzing samples from 8 newborns with congenital CMV infection treated with 15 mg/kg of Valganciclovir (GCV prodrug) twice daily.

Conclusion

The analytical method developed and validated is rapid and controllable in quality simple. The method showed high resolution, high sensitivity, good linearity, precision, accuracy and sensitivity. The inter-day variability is very low, and samples are stable in the condition tested up to 7 days. The recovery of GCV gave rise to a satisfactory recovery, thus avoiding the use of internal standards. Comparing our method to other proposed in the literature, it determines the GCV in smaller plasma concentrations, and is less expensive as it use a smaller variety of solvents. Results obtained in this study demonstrate its usefulness for therapeutic monitoring.

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