

PQ

for DECADE Elite, Lite, II I and Intro

171.0023P, Edition 18, 2026



Copyright ©2026, Antec, The Netherlands. Contents of this publication may not be reproduced in any form or by any means (including electronic storage and retrieval or translation into a foreign language) without prior agreement and written consent from the copyright of the owner. The information contained in this document is subject to change without notice.

ROXY, ALEXYS, DECADE, DECADE Elite, DECADE Lite, DECADE II, INTRO, Flexcell, SenCell, μ -PrepCell, ReactorCell, SynthesisCell, ISAAC, HyREF are trademarks of Antec. Whatman™ (word and device) and Whatman™ (word only) are trademarks of Whatman International Ltd. SOLVENT IFD™ and AQUEOUS IFD™ are trademarks of Arbor Technologies, Inc. Clarity®, DataApex® are trademarks of DataApex Ltd. Microsoft® and Windows™ are trademarks of Microsoft Corporation. Excel is a registered trademark of the Microsoft Corporation.

The software and the information provided herein is believed to be reliable. Antec shall not be liable for errors contained herein or for incidental or consequential damages in connection with the furnishing, performance, or use of software or this manual. All use of the software shall be entirely at the user's own risk.

Table of contents

Chapter 1	3
Introduction	3
Supported configurations	3
Chapter 2	5
PQ test.....	5
Principle	5
Required parts and consumables	6
Preparations	8
HPLC-ECD test conditions for PQ	10
PQ procedure	11
Processing the PQ results	12
Chapter 3	15
Specifications.....	15
PQ test on gold electrodes	15
PQ test on glassy carbon electrodes	15
Chapter 4	17
OQ-PQ results summary	17
Chapter 5	19
OQ-PQ results summary of optional 2 nd cell	19
Chapter 6	21
PQ certification	21
Engineer	21
Reviewer/customer	21
Instrument	22
Test materials	23
Test devices	23
Comments	24
Chapter 7	25
Non-conformance record.....	25

CHAPTER 1

Introduction

This document describes the Qualification procedure as advised by the manufacturer. It is a result from our interpretation of many regulations and laboratory practices. In addition, feedback from users and representatives helped us to finalize this procedure.

In this document, all qualification checks have to be approved or should be marked "n.a." if not applicable. Any deviation observed must be documented in the 'non-conformance' record. All relevant documents regarding this operational qualification must be filed together in one location.

As regulations and customer requirements may change, the manufacturer reserves the right to introduce changes without prior notice. For details on functionality, operation and theory refer to the instrument user manuals.

Supported configurations

The PQ procedure in this document is applicable to an HPLC-ECD system with

- an Antec electrochemical detector
 - DECADE Elite™
 - DECADE Lite
 - DECADE II™
 - DECADE™
 - INTRO™

- with an Antec flow cell with one of these working electrode materials
 - **glassy carbon** (GC)
 - **gold** (Au)

Flow cells with working electrodes other than Au or GC are not supported with PQ procedures.

CHAPTER 2

PQ test

Principle

For testing the performance of the HPLC/ECD system, a test substance is injected onto a Flow Injection Analysis (FIA) tubing-assembly, and repeatability, linearity & peak height is measured.

Different detection modes for GC and Au working electrodes

In case of the use of a flow cell with glassy carbon electrode, the detection method is set to direct current (DC) mode, which will apply one working potential. In case of testing with a gold working electrode, Pulsed Amperometric Detection (PAD) mode is used.

Test parameters

Important parameters to characterize and check the performance of all relevant parts of a detection system are:

- Repeatability
- Linearity
- Detector response
- Background current (only with GC electrode)
- Signal-to-noise ratio (only with GC electrode)

Not only the detector and flow cell performance but also those of pump and autosampler are taken into account with these parameters. In case of testing the performance with a gold flow cell, the background current, noise and S/N measurement are not relevant, as in PAD mode the signal is digitally processed.

As the detector response is affected by electrode size and injection volume, different values are specified for different combinations of hardware.



Figure 1. PQ tubing kit installed in detector oven and connected to flow cell.

Required parts and consumables

To perform the PQ, a specific PQ consumables kit is necessary (Table 1). It consists of the documents, chemicals and a PQ FIA tubing set (consisting of a restriction and mixer tubing and parts to connect to 1/16" and 1/32" receiving ports). During PQ, the PQ FIA-coil must be installed in place of the column (see Figure 2).

Two different PQ consumables kits are available:

- pn. 250.3048A, specific for testing with a glassy carbon (GC) electrode
- pn. 250.3046A, specific for testing with a gold (Au) working electrode

Table 1. Two versions of PQ consumables kits

Part	Antec pn.	Qty
PQ consumables incl FIA-coil, ECD in DC mode		
PQ FIA tubing set for Flow Injection Analysis	250.1052	1
Concentrated buffer for PQ	250.1064	1
MOPEG 4.0 µmol/L, 8 mL for PQ	250.1062A	1
Document: PQ for D2 Elite Lite and ROXY	171.0023P	1
Document: OQ for D2 Elite Lite and ROXY	171.0023O	1
PQ consumables incl FIA-coil, ECD on Au at pH 13		
PQ FIA tubing set for Flow Injection Analysis	250.1052	1
Glucose for PQ (dry powder; for 3 mL stock solution of 20 mM)	250.1067	1
Document: OQ for D2 Elite Lite and ROXY	171.0023O	1
Document: PQ for D2 Elite Lite and ROXY	171.0023P	1

Additional consumables and chemicals that are necessary to perform the PQ procedure are:

- Autosampler vials
- 1 L ultra-pure water from a water purification system (with resistivity > 18 MOhm.cm and TOC < 10ppb)

For test in DC mode on **GC** electrodes:

- 50 mL methanol, HPLC-grade or better

For test in PAD mode on **Au** electrodes:

- 50% sodium hydroxide solution, carbonate free HPLC grade (commercially available solution)

Configuration for systems with a post-column addition pump

When testing an HPLC system with a post-column addition pump (for example in case of analysis of certain aminoglycosides), the PQ FIA tubing must be connected upfront the post-column mixer (see Figure 2).

The flow rate ratio of the main and post column pump should be set to a ratio of 2:1, for example 1 mL/min (main) and 0.5 mL/min (post column addition). The final NaOH concentration **after mixing** must be as specified in Table 2. With the main pump delivering water, the post-column pump must therefore deliver a 3x higher concentration of NaOH to make up for the dilution. Thus, use a solution of **300 mmol/L NaOH for the post column pump**. The performance specifications for these conditions are the same as described for a system with only 1 pump. Make sure to prepare the standard in water, which is the mobile phase in this 2-pump set-up.

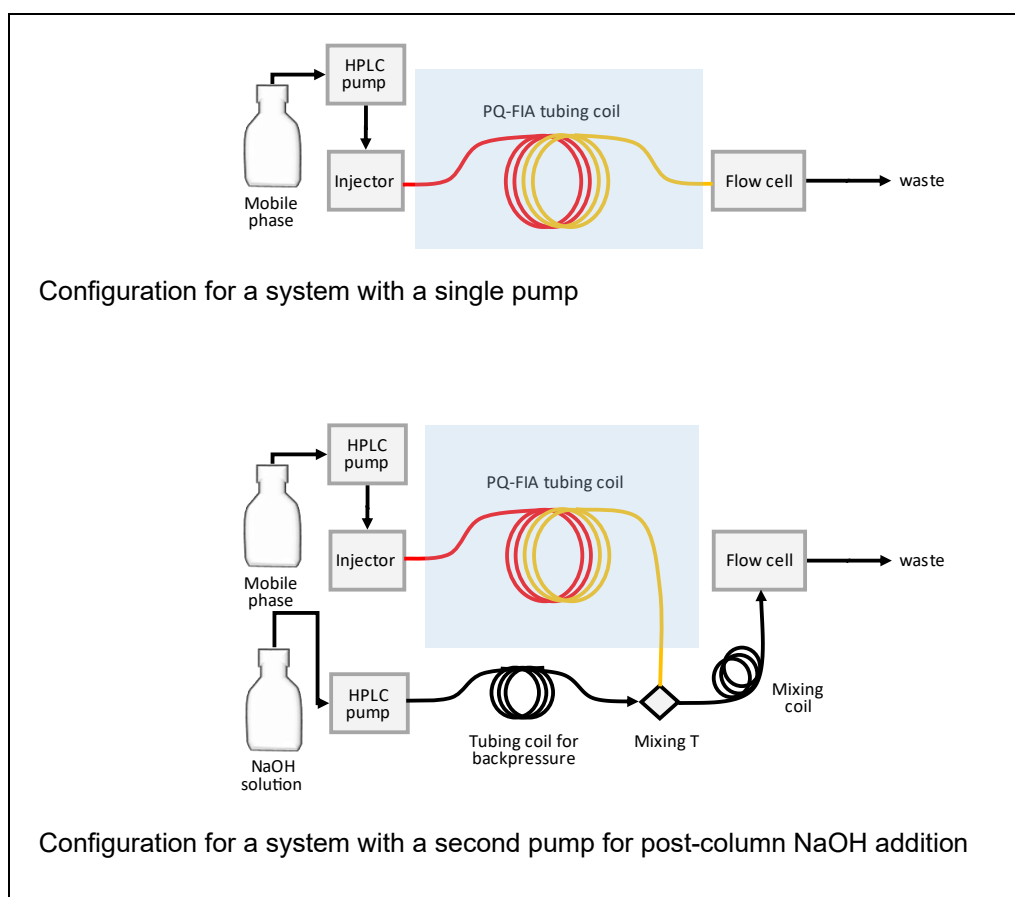


Figure 2. For PQ the analytical column is replaced by the PQ-FIA tubing coil (assembly of 3 m PEEK tubing with ID of 125 μm and 10 m PEEK tubing with ID of 500 μm).

Preparations

For a successful performance test, it is important that the HPLC-EC system has been optimized and is in top condition.

OQ

1. Perform the Operation Qualification procedure (pn. 171.0023O) before performing the PQ test. Results of the OQ are to be summarized in the PQ results table on page 17.

Flush the system

2. Prepare a bottle with degassed water and use it to flush the system up to the injector.
3. Connect the red-striped end of the PQ FIA tubing directly to the valve. If needed use the 1/32"OD tubing adapter.
4. Flush the FIA tubing for 10 min with water at 1 mL/min.
5. Prepare the mobile phase according to Table 2.
6. Connect the mobile phase, purge the pump(s) and run about 30 mL through the system.

Install the flow cell and let the signal stabilize

7. Clean the flow cell (see user manual) and assemble with a 50 µm spacer, AST 2 setting.
8. Connect the flow cell using the dedicated flow cell connectors (see Figure 2) and ensure that the flow cell is filled air-free with mobile phase.
9. Let the mobile phase flow through the cell for 5 min.
10. Set the detector conditions as given in Table 2 and turn on the cell.



Do not turn the cell on when mobile phase has not yet completely filled the cell. Turning on a cell that still has air or water inside can damage the cell!

11. Wait for the system to stabilize for at least 30 min before starting the PQ HPLC tests.

Prepare the sample

12. For testing flow cells with glassy carbon electrode only: the MOPEG solution is a ready to use standard and only needs to be diluted in case the linearity check is done with diluted standard instead of injection volume variation.
13. For testing flow cells with gold electrode only: Glucose needs to be dissolved and diluted:
 - Add 3 mL of mobile phase to the vial with glucose and shake to dissolve thus making a 20 mM stock solution. Note that for systems running with a post-column pump, the 'mobile phase' is water not NaOH solution.

- Dilute the 20 mM stock solution 1000x with MP. For example, dilute in 3 consecutive steps: mark 3 empty vials clearly as "1", "2", "3" and add 900 μL mobile phase to each of the 3 vials. Add 100 μL stock solution in vial 1, close and shake well. With a new clean pipet tip take 100 μL solution from vial 1 and add to vial 2, close and shake. With a new clean pipet tip take 100 μL solution from vial 2 and add to vial 3. Vial 3 now contains 1 mL of 20 $\mu\text{mol/L}$ glucose standard ready for a PQ test injection.

For cells with gold working electrode the PQ standard solution is 20 $\mu\text{mol/L}$ glucose in mobile phase (MP). Dissolve and dilute the glucose powder in mobile phase (100 mM NaOH or water in case of post-column addition) prior to use.

HPLC-ECD test conditions for PQ

Table 2. HPLC-EC conditions for PQ.

General settings	
PQ hardware	PQ FIA tubing set (250.1052) installed in place of the column
Flow rate (signal)	1 mL/min (up to 1.5 mL/min is allowed)
Temperature	35 °C (both flow cell and PQ FIA tubing inside the detector oven)
Integrator	Data acquisition frequency: < 5 Hz
Flow cell spacing	SenCell: AST setting '2' VT-03 and FlexCell with black PTFE inlet block: 50 µm spacer FlexCell with stainless steel inlet block: 130 µm spacer
Settings for testing with <u>glassy carbon</u> working electrodes	
Mobile phase	1 bottle concentrated PQ buffer (pn. 250.1064), 50 mL MeOH*, mixed with water* to a final total volume of 1000 mL. Note: PQ buffer already contains chloride for use with an ISAAC reference electrode.
Sample	MOPEG 4 µmol/L (pn. 250.1062A)
E cell	+610 mV (vs HyREF); +800 mV (vs sb); +650 mV (vs ISAAC)
Filter	0.1 Hz
Range (for signal)	200 nA/V for 2 mm, 3 mm and FlexCell GC electrodes 10 nA/V for 0.7 mm GC electrodes
Range (for noise)	1 nA/V
Flow rate (for noise)	1 mL/min (0 mL/min is allowed as alternative for the normal flow rate)
Settings for testing with <u>gold</u> working electrodes	
1 pump set-up	<i>Mobile phase</i> : 100 mM NaOH* in water*
Solutions and flow rate for post-column addition set-up with 2 pumps	<i>Mobile phase</i> : 1 mL/min water* <i>Post-column solution addition</i> : 0.5 mL/min 300 mM NaOH* in water* Note: the final concentration after mixing is then 100 mM NaOH
Sample	20 µmol/L glucose prepared in <i>mobile phase</i>
E (PAD mode) vs. HyREF or Ag/AgCl sb	For DECADE Elite: E1, E2, E3, E4 = 100, -2000, 600, -100 mV (E5 0 mV); t1, t2, t3, t4, ts = 400, 20, 10, 70, 200 ms (ts: 0 ms) For DECADE II: E1, E2, E3 = 100, -2000, -100 mV t1, t2, t3, ts = 400, 20, 80, 200 ms For FlexCell with stainless steel inlet block: E1, E2, E3: +0.05, +0.75, -0.15 V; t1, t2, t3: 0.4, 0.15, 0.45 s (ts: 300 ms)
Icell	About 0.5 - 2 µA
Range & filter	1 µA/V, off
Specific settings for DECADE I and INTRO	
Filter	Signal: 0.1 s; Noise: 1 s
Noise analysis	DECADE: pA range; INTRO: att x100 = off

* Ensure that all chemicals meet the minimum purity requirements as specified on page 6.

PQ procedure

Test injection

Run a test chromatogram with the test solution. Check the test-chromatogram peak time and optimize the automated integration parameters if necessary.

In case of using a 20 or 100 μL sample loop on the injector: adjust the detector range if the test peak height is larger than 50% of the full-scale range: the analyses for linearity require a larger injection volume and the signal may run off scale if the range setting is set too low.

Linearity test

A 5-point equidistant calibration plot has to be set up for the linearity evaluation. Data can be generated by varying the concentration or injection volume:

- Varying injection volumes: inject 5 different volumes of the test solution, for example
 - * for injectors with a 20 μL loop: 4, 5, 6, 7, 8 μL
 - * for injectors with a 100 μL loop: 20, 25, 30, 35, and 40 μL
- Varying concentrations: dilute the test solution by hand thus generating equidistant concentrations, for example
 - * 0.8, 1.6, 2.4, 3.2 and 4.0 $\mu\text{mol/L}$ in case of MOPEG
 - * 4, 8, 12, 16, 20 $\mu\text{mol/L}$ in case of glucose

The most preferred method is varying the injection volume because it eliminates manual dilution errors.

Repeatability test

Repeatability is evaluated based on 8 subsequent analyses of the test solution.

Noise measurement and background current (for GC electrodes only)

Check the cell current (**I cell**) at the detector (e.g. read from the display) and write it in the Results Summary table (page 17).

Record a 15 min baseline trace using the setting specified in Table 2. When the pump is suspected to contribute to the noise level, the pump may be switched off or set to another optimum flow rate.

Processing the PQ results

The PQ procedure results in linearity data, 8 repeatability chromatograms, and a baseline trace (for GC electrodes only). See Fig. 3 for example chromatograms.

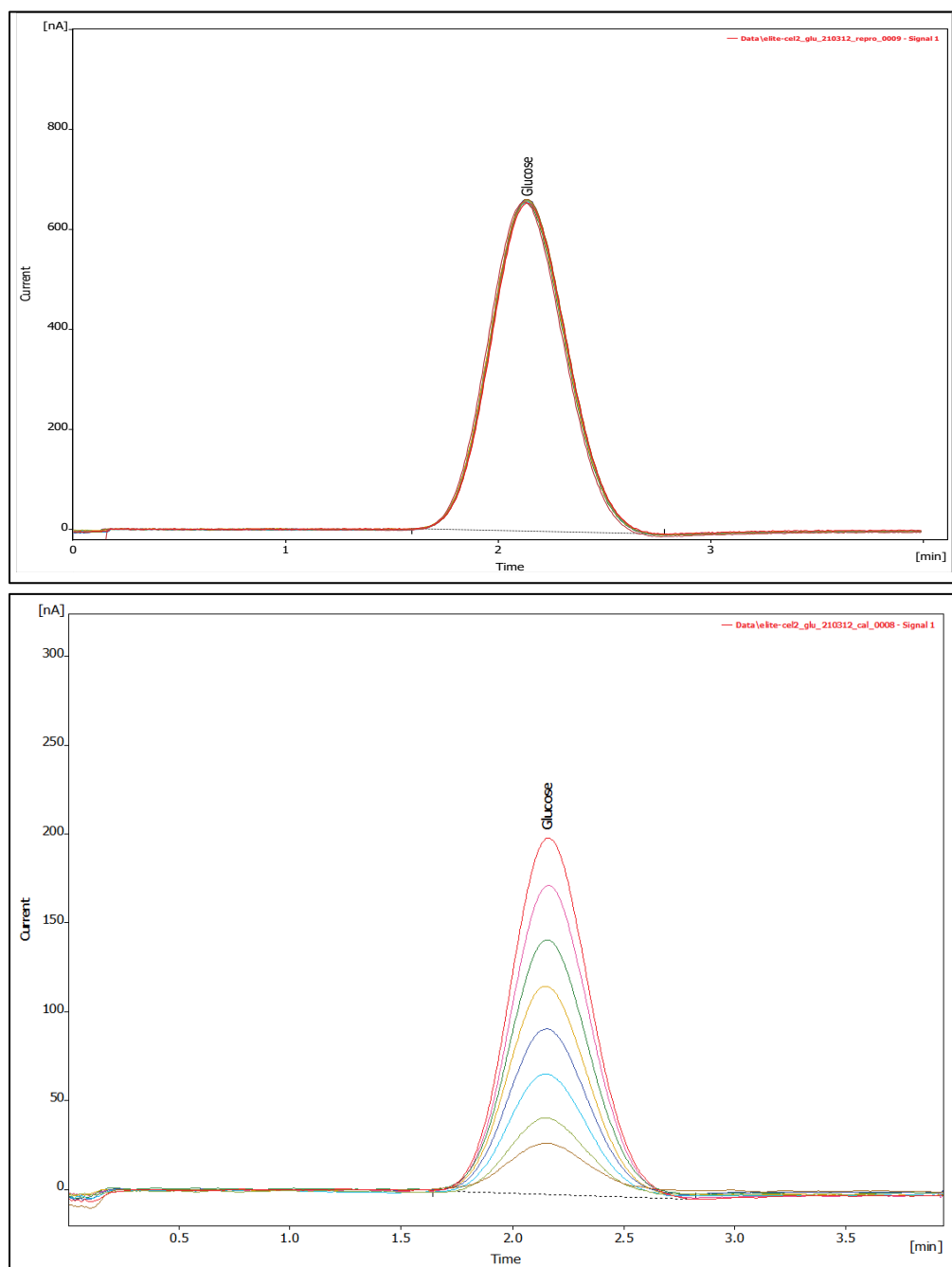


Fig. 1. Example chromatograms for 20 μM glucose solution prepared in 100 mM NaOH (mobile phase); top (repeatability test): overlay of 8 repetitive injections of 20 μL (full loop); bottom (linearity test): overlay of 8 injections of 1, 2, 3, 4, 5, 6, 7 and 8 μL (partial loopfill).

1. Select the 8 'repeatability' chromatograms and create a report containing
 - retention time
 - peak heights
 - peak areas
2. Calculate the average values and the relative standard deviation as percentage of the average (%RSD).
3. Process the linearity data and calculate the correlation coefficient between peak area and concentration or injection volume.

Write the results down in the PQ results table on page 17.

Contact your supplier if the final result of the HPLC PQ procedure is 'failed'. Note also that a 'failed' PQ is not necessarily related to the detector. For example, repeatability failures are often an HPLC problem (autosampler).

Additional parameters for GC electrodes:

4. Open the baseline trace and evaluate the noise as described in the OQ document (171.0023O), or use any other software that is capable of the calculating the baseline peak-to-peak noise level (ASTM).
5. Calculate the S/N ratio (=peak height/noise level) after obtaining the noise value. The average peak height from the repro-evaluation is used as "signal".

Write the results down in the PQ results table on page 17.



To be able to reuse (part of) a PQ tubing set, flush it with water before storage.

Before flushing the system, SWITCH OFF AND REMOVE THE CELL!

Recalculation to the correct unit - signal from mV to nA

In case the detector is connected to an AD convertor, this will normally generate a signal with the units in mV (or μV). Use the detector range setting to convert these values to the actual ECD *current*.

$$\text{Range (nA/V)} * \text{Signal (mV)} = \text{Signal (pA)}$$

For example, at the 200 nA/V range, a signal of 300 mV is actually a current of:

$$200 \text{ nA/V} * 300 \text{ mV} = 6000 \text{ pA} (= 6 \text{ nA})$$



Use the correct units when multiplying the range and analog output signal!

CHAPTER 3

Specifications

PQ test on **gold** electrodesPeak height specification of 20 μM glucose test solution

Injection volume	Au working electrode				
	0.7 mm*	2 mm*	3 mm*	FlexCell**	
				50 μm	130 μm
5 μL	> 1.3 nA	> 6.3 nA	> 13 nA	> 25 nA	> 15 nA
20 μL	> 5 nA	> 25 nA	> 50 nA	> 100 nA	> 55 nA

*AU WE: 0.7 mm is $\mu\text{VT-03}$, 3 mm is VT-03, 2 mm is SenCell

**FlexCell: only inlet block stainless steel comes with 130 μm spacer thickness (black PEEK has 50 μm)

PQ test on **glassy carbon** electrodesPeak height specification of 4 μM MOPEG test solution

Injection volume	GC working electrode			
	0.7 mm*	2 mm*	3 mm*	FlexCell
2 μL	> 0.1 nA	> 0.5 nA	> 1 nA	> 1 nA
5 μL	> 0.2 nA	> 1 nA	> 2 nA	> 2.5 nA
10 μL	> 0.4 nA	> 2 nA	> 4 nA	> 5 nA
20 μL	> 0.8 nA	> 4 nA	> 8 nA	> 10 nA

*GC WE: 0.7mm is μVT03 , 3mm (2.7mm) is VT03 sn 4xxxx 2mm is SenCell or VT03 sn 2xxxx.

SN-ratio specification for 4 μM MOPEG test solution

Injection volume	GC working electrode			
	0.7 mm	2 mm	3 mm	FlexCell
2 μL	> 500	> 500	> 500	> 250
5 μL	> 1000	> 1000	> 1000	> 500
10 μL	> 2000	> 2000	> 2000	> 100
20 μL	> 4000	> 4000	> 4000	> 2000

Noise and background current specification

	GC working electrode			
	0.7 mm	2 mm	3 mm	FlexCell
I cell	< 3 nA	< 8 nA	< 12 nA	< 20 nA
noise p-p	< 2 pA	< 6 pA	< 8 pA	< 14 pA

Dummy cell test - noise

The noise spec for a dummy cell test with a ROXY: noise < 4.0 pA. For all other devices: noise < 2.0 pA.

CHAPTER 4

OQ-PQ results summary

Injected volume of sample for response and repeatability tests: μL

Electronic test results and HPLC test results

	Specified* ¹	Measured	Result* ²
<u>ELECTRONIC TESTS</u> * ³			
Dummy cell test			
Current (I-cell)	2.67 \pm 0.05 nA nA
Noise p-p	< pA pA
Analog output test			
Output at 5 nA/V	530 \pm 10 mV mV
<u>HPLC TESTS</u>			
Chromatogram			
Peak time	< 3 min min
Response			
Height	> nA nA
Repeatability			
%RSD t	< 0.5 % %
%RSD area	< 3.0 % %
Linearity			
Correlation coefficient r	> 0.997
Tests for GC WE only * ⁴			
I-cell	< nA nA
Noise	< pA pA
S/N ratio	>

*¹ Specifications for some of the HPLC tests are hardware dependent: check the applicable specs on page 15 and copy the specified value into this table.

*² Fill in 'passed' or 'failed'

*³ for 'electronic test' results, copy the measurement values from the OQ document.

*⁴ Tests for background current and sensitivity are applicable only in case of using a glassy carbon working electrode in the flow cell. When testing with a **gold** working electrode, fill in n.a.

Final result*² _____

Verified by (customer):

Deviations (Y/N):

Comments:

CHAPTER 5

OQ-PQ results summary of optional 2nd cell

This page is for use on DCC detectors that are tested in parallel or serial configurations, otherwise strike through with n.a. (not applicable)

Injected volume of sample for response and repeatability tests: μL

Electronic test results and HPLC test results on 'Cell 2'

	Specified ^{*1}	Measured	Result ^{*2}
<u>ELECTRONIC TESTS</u>^{*3}			
Dummy cell test			
Current (I-cell)	2.67 \pm 0.05 nA nA
Noise p-p	< pA pA
Analog output test			
Output at 5 nA/V	530 \pm 10 mV mV
<u>HPLC TESTS</u>			
Chromatogram			
Peak time	< 3 min min
Response			
Height	> nA nA
Repeatability			
%RSD t	< 0.5 % %
%RSD area	< 3.0 % %
Linearity			
Correlation coefficient r	> 0.997
Tests for GC WE only^{*4}			
I-cell	< nA nA
Noise	< pA pA
S/N ratio	>

*1 Specifications for some of the HPLC tests are hardware dependent: check the applicable specs on page 15 and copy the specified value into this table.

*2 Fill in 'passed' or 'failed'

*3 for 'electronic test' results copy the measurement values from the OQ test.

*4 Tests for background current and sensitivity are applicable only in case of using a glassy carbon working electrode in the flow cell. When testing with a **gold** working electrode, fill in n.a.

Final result^{*2} _____

Verified by (customer):

Deviations (Y/N):

Comments:

C H A P T E R 6

PQ certification

The Performance Qualification has been carried out in accordance to the PQ procedure and to the satisfaction of both parties.
All tests as described in this document have been completed, and all results are within specifications, or clearly indicated if not.

Engineer

The undersigned engineer certifies that he/she is trained and qualified to perform a PQ on Antec devices.

Name
Initials
Company

.....
Date Signature

Reviewer/customer

The undersigned reviewer/customer is authorized to sign and accepts that the above-mentioned engineer is trained and qualified to perform a PQ on Antec devices.

Reviewer/Customer
Initials
Job title
Company & Dept.

.....
Date Signature

LC-EC instrument

EC detector model*

- DECADE II
- DECADE Elite
- ROXY Exceed
- ...

p/n: s/n:

Firmware version

Boot version

Instrument has DCC option (Y/N)

Grounding kit option installed (Y/N)

Flow cell type

p/n: s/n:

Working electrode type:

Reference electrode type:

Flow cell type (2nd)**

p/n: s/n:

Working electrode type:

Reference electrode type:

Autosampler /injector s/n:

Installed sample loop size: μL

Degasser model s/n:

Pump 1 model s/n:

Pump 2 model*** s/n:

* tick the right model

** enter the info of a second flow cell only in case of use in a DCC detector.

*** enter the info of a second pump in case post column addition or parallel set-up with DCC detectors was used.

Verified by (customer): Deviations (Y/N):

Comments:

Chromatographic Data-acquisition Software (CDS)

CDS package name

Version and revision nr.

.....

.....

Data acquisition details

EC detector is connected to an AD convertor (Y/N)

EC detector is placed under software control (Y/N)

If 'Y': **DECADE Elite driver version and revision nr.**

Test materials

Standard*

Lot nr Exp. date

Mobile phase

* Fill in 'glucose' or 'MOPEG'

Test devices

Dummy cell* p/n: 250.0040 s/n:

Voltmeter or AD signal s/n:

*s/n: entering more than one s/n is allowed for DCC detectors.

Other relevant information

Description	

Verified by (customer):

Deviations (Y/N):

Comments:

Comments

Verified by (customer):

Deviations (Y/N):

Comments:

C H A P T E R 7

Non-conformance record

Any case of non-conformance found during the qualification procedure should be documented and signed for acceptance or corrective action taken.

Ref.	Non-conformance and action taken or acceptance	Signature customer	Sign. executing technician
1	
2	
3	
4	
5	
6	

Verified by (customer):