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Level IV Data Package

MWH Group 212263

Method: 7196 CRVI

Sample No.:

2708030231

2708030232

2708030233

2708030234

2708030235

SM 7196A QC Check List	Cr-VI	
Analyst WBY	Analysis date \$\frac{\$\frac{3}{3}}{2}\frac{3}{2}\text{Reviewer/Date} \tag{UV}	8/6/07
Instrumnet: HACH DR/4000V		
,All sample analyzed	within 24 Hrs. holding time	
	centraction below the high standard or linear range of this batch erranged samples marked for dilution and rerun	
ALL QC	tion verifications within +/- 10%	
	D within +/+ 15%	
	within +/-30%	
MRL within	** ***	
	k <1/2 MRL	
*	reen MS/MSD is within +/-/20%	
All pH of ti	he samples are 7	
No more than 20 sar	mples per batch	
MS is run at frequen	cy of 1 every 10 samples and MSD is	
run at freq	quency of 1 every 20 samples	
QIR needed for failed	d QC	
Special Det Code no	oted on the cover sheet	

instrument: Hach DR 400	ov			Cr-VI b	EPA I	Method	7196A							
Analyst: Walto	s Há	/		79			Page /.of		Start time:	11	:15	End time:	12:00	$, \neg$
Cal. Stock Std:	Use Built-In Curve	•		Cal. Stock Std exp. Date: 08/10/2007				Cal. Worki	ng Stat:		070511-1	/		125
LCS Stock Sta:	HACH DR4000V			LCS Stock S	Std exp. Da	te: 08/10/20	07 /	LCS Work	ng Std:		070511-2	/		\exists
Correlation Coeff:				Slope:			Y-intercept:			7	True Value		0,05PPM /	7
Sample		Di	ution		Tur	t Blk		-				***************************************	·····	\neg
		sample	total		(same	dilution)	Calc.	Reported	Sampling	piH		Comr	nente	
		mts	mls	abs	abs	res abs	mg/L	mg/L	date		Anal. Time	T		_
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Std 1 (0.005 ppm)		10	10	0,010			0.525	0.505		T		(33)	\	\neg
Std 2 (0.020 ppm)		10	10	0038		***************************************	นจระ	ر ١٤٥٠	7	1		(10 3)	5	[
Std 3 (0.050 ppm)		10	10	0,092	·		005.	0250		T		(00)	······································	
Std 4 (0.290 ppm)		10	10	:360			0 197	0.197.	/	T		(385		
Std 5 (0.500 ppm)		10	10	U ÝŸ			0.483	1.483		T	(9667		一
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LCS - 1 (0.05 ppm)		10	10	0376			0052	0052	//			(su	Limit: 85 - 1159	
Sample Log #	イルーリ Sample ID	5	10	26/2			4.338	4016		7	13:52		UR	$^{\sim}$
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LCS - 2 (0.05 ppm)		10.	1	093				025		十	·		limb of acce	-
0.020 ppm/CCV		10		6.036				0.020		\dashv		23/	Limit: 85 - 115%	\dashv
Blank/CCB		10		0.300				237	7	\dashv			Limit: 90-110%	\dashv
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	v 115)			8/6/07					color Reagent EXP, Date	. MAC	in Perma Cr	em Chrom	a ver3	\dashv

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all 4sxs were sampled at 10:00. Fold to. Arrived past. 10:00 the wext day.

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Standard
Preparation
Worksheet
&
Certificate of
Analysis

5 ppm Crtb Std					
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	MW #	#: WHO70206			
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WBM	Amoun	t: 100 nl			
Rosm Temp	Lot #:				
Comment	Standard	Concentration			
* ME OCO ***		5ppm			
t == (-16 0)					
3 fph ar ac	MW #:	W1070206-			
216 1071 1 1	By:	wsn_			
516 1071 1 1	Matrix:	AQ KK			
Wyn	Amount:	100m/			
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Comment	Standard	Concentration			
R 201081		5ppm			

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5 / 1671	IVIVV #:	WH070511-			
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1/8n	_				
MSh Amount: 100n					
even leng	LOT#:				
Comment	Standard	Concentration			
R 201632		5ppm			
R Z0/632		5ppm			
R 20/632		5ppm			
R Z0/632		5ppm			
R 20/632		5ppm			
R Z 0 / 632		5ppm			
	Comment PME 060403 5 ppn Cr16 QC 2 16 107 1 1 1 W37 Rown Temp Comment R 201081	Comment Standard PME0604003			

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	Reagent Preparation Documentation		Page: 11
Reagent:	Co16 QCStd. 5111107111 811016807111	RANA	#: WHO70511
Date Received/Prepped:	5/11/07/		#
Date Expired:	81/01-011	. Matr	iv Ok 1170
Manufacturer:	WRH	Amou	ix: 100m/
Storage Condition:	Room Temp	Lot	#:
Component	Comment	Standard	Concentration
Cx 16 0.5ml	Rt 201090		5 ppn
Comment:			
Reagent: Date Received/Prepped:	5 Mm Cr to Cal Std \$191071 1 1 11191071 1 1 FAJ Room Foo	MW i	#: fg 080907 - 1: fg 1: agons 12 H 1: jormL
Date Expired:	11/9/07/	Matrix	C: ccseous 12 H
/lanufacturer:	FAI	Amoun	: formL
Storage Condition:	Room temp	Lot #	:
Component	Comment	Standard	Concentration
Cr+4 0.5 ml	R 201090	~	5 ppm
	•		
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omment.			
leagent:	Crts GC std	MW#	: Fgi 080907-
ate Received/Prepped:	819107111	Ву	Fg 080907-
ate Expired:	1/19107111		agens 12 H,
anufacturer:	PAJ		100 mL
orage Condition:	Room temp	Lot#:	
Component	Comment	Standard	Concentration
Cr 46 0.5 mL	R 201632		5 ppm

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Maddecate

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QUARTER CAS

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Service S



CERTIFICATE OF ANALYSIS

195 Lehigh Avenue, Suite 4 Lakewood, New Jersey 08701 - USA inorganicventures.com

tei: 800.669.6799 + 732.901.1900 fax: 732.901.1903

into@inorganicventures.com

1.0 INORGANIC VENTURES is an ISO Guide 34:2000 registered Certified Reference Material (CRM) Manufacturer (Certificate #883-02). The certificate is designed and the data is determined in accordance with ISO Guide 31:2000 (Reference Materials-Contents of Certificates and Labels), ISO Guide 34:2000 "Quality System Guidelines for the Production of Reference Materials," and ISO Guide 35:1989 "Certification of Reference Materials - General and Statistical Principals."

2.0 DESCRIPTION OF CRM 1000 μg/mL Chromium (+6) in H20

R# 201632

Catalog Number:

CGCR(6)1-1, CGCR(6)1-2, and CGCR(6)1-5

Lot Number:

Z-CR02152

Starting Material:

(NH4)2Cr2O7

Starting Material Purity (%):

99.989259

Starting Material Lot No:

F04N14

Matrix:

H20

3.0 CERTIFIED VALUES AND UNCERTAINTIES

Certified Concentration:

 $1000 \pm 3 \mu g/mL$

Certified Density:

0.999 g/mL (measured at 22° C)

The Certified Value is based upon the most precise method used to analyze this CRM. The following equations are used in the calculation of the certified value and the uncertainty:

Certified Value $(\bar{x}) = \Sigma \underline{x}_i$

(x̄) = mean

x_i = individual results

n = number of measurements

Uncertainty (±) = $2[(\Sigma s_i)^2]^{1/2}$ (n) $(r)^{1/2}$

 Σ s_i = The summation of all significant estimated errors

(Most common are the errors from instrumental measurement, weighing, dilution to volume, and the fixed error reported on

the NIST SRM certificate of analysis.)

The independent samples t-test was used to determine if there is agreement between the above assay methods at the 95% confidence interval. Both methods were compared and showed agreement within the stated uncertainties. This agreement is a confirmation of the accuracy of this CRM.

4.0 TRACEABILITY TO NIST AND VALUES OBTAINED BY INDEPENDENT METHODS

- "Property of the result of a measurement or the value of a standard whereby it can be related to stated references, usually national or international standards, through an unbroken chain of comparisons all having stated uncertainties." (ISO VIM, 2nd ed., 1993, definition 6.10)
- This product is Traceable to NIST via an unbroken chain of comparisons. The uncertainties for each certified value are reported, taking into account the SRM uncertainty error and the measurement, weighing and volume dilution errors. In rare cases where no NIST SRMs are available, the term 'in-house std.' is specified.

4.1 Assay Method #1 1000 ± 3 μg/mL

Redox NIST SRM 136e Lot Number: 980702

Assay Method #2 1001 ± 4 µg/mL

ICP Assay NIST SRM 3112a Lot Number: 990607

- BALANCE CALIBRATION All balances are checked daily using an in-house procedure. The weights used for testing are annually compared to master weights and are traceable to the National Institute of Standards and Technology (NIST). The NIST Traceability numbers are 692476. - Class 1 and 692476A - Class 2. The NIST test number is 822/260017-98. All analytical balances are calibrated every 4 months. The balances are calibrated with a class 1 and/or class 2 analytical weight set. These weights are tested annually by a NIST / NVLAP accredited calibration lab. The NIST test number is 822/260017-98.
- THERMOMETER CALIBRATION The thermometers used in the determination of the final densities are calibrated vs standard. thermometer No. 903-2680 which was certified in accordance with the procedures outlined by ASTM E77-87 and NIST Monograph 150 using NIST Test Nos. and Std Nos.: 769543, 217368/769543, 217368/P14452, 176240/P14452, 176240. Thermometers which are not calibrated vs standard thermometer No. 903-2680 are traceable to NIST Identification Nos. 92564, 119016, 471047 and NIST test report Nos. 811/258522, 811/2557078, and 236090.
- GLASSWARE CALIBRATION An in-house procedure is used to calibrate all Class A glassware used in the manufacturing and quality control of CRM's.

TRACE METALLIC IMPURITIES (TMI) DETERMINED BY ICP/MS AND ICP-OES IN µg/mL 5.0

CRM's solutions are tested for trace metallic impurities by Axial ICP-OES and ICP-MS. The result from the most sensitive method for each element, is reported below. Solutions tested by ICP-MS were analyzed in an ULPA-Filtered Clean Room. An ULPA-Filter is 99.9985% efficient for the removal of particles down to 0.3 µm.

			14 D	LL To
<u>Q</u> Al < 0.00090	M Dy < 0.01185	<u>Q</u> Li < 0.00002	<u>M</u> Pr < 0.00059	<u>M</u> Te < 0.05923
<u>M</u> Sb < 0.00099	M Er < 0.00987	M Lu < 0.00079	M Re < 0.00197	<u>M</u> Tb < 0.00059
<u>M</u> As < 0.01974	<u>M</u> Eu < 0.00592	<u>Q</u> Mg < 0.00030	M Rh < 0.00197	<u>M</u> TI < 0.00197
<u>M</u> Ba < 0.01974	<u>M</u> Gd < 0.00197	<u>M</u> Mn < _{0.00790}	M Rb < 0.00197	<u>M</u> Th < 0.00197
<u>M</u> Be < 0.00099	<u>M</u> Ga < 0.00197	<u>O</u> Hg < 0.01500	<u>M</u> Ru < 0.00395	\underline{M} Tm < 0.00079
<u>M</u> Bi < 0.00079	<u>M</u> Ge < 0.01185	<u>M</u> Mo < 0.00395	<u>M</u> Sm < 0.00197	<u>M</u> Sn < 0.00987
<u>Q</u> B < 0.01000	<u>M</u> Au < 0.00592	M Nd < 0.00395	M Sc < 0.01974	<u>Q</u> Ti < 0.00100
<u>M</u> Cd < 0.00592	<u>M</u> Hf < 0.00395	<u>M</u> Ni < 0.01579	M Se < 0.01579	<u>M</u> W < 0.01974
<u>O</u> Ca _{0.00027}	<u>M</u> Ho < 0.00099	<u>M</u> Nb < 0.00099	<u>Q</u> Si < 0.20000	<u>M</u> U < 0.00395
M Ce < 0.00987	<u>M</u> in < 0.01974	n Os	M Ag < 0.00395	Q V - < 0.02000
M Cs < 0.00059	<u>M</u> Ir < 0.00987	M Pd < 0.00987	Q Na 0.00300	M Yb < 0.00197
<u>s</u> Cr	<u>O</u> Fe < 0.01000	<u>O</u> P < 0.04000	<u>M</u> Sr < 0.00099	<u>M</u> Y < 0.07897
M Co < 0.00592	<u>M</u> La < 0.00099	M Pt < 0.00395	į S	\underline{O} Zn < 0.00400
<u>M</u> Cu < 0.01185	<u>M</u> Pb < 0.00592	<u>O</u> K 0.32485	M Ta < 0.01382	<u>M</u> Zr < 0.00987

M - Checked by ICP-MS

O - Checked by ICP-OES i - Spectral Interference

n - Not Checked For

s - Solution Standard Element

INTENDED USE 6.0

For the calibration of analytical instruments including but not limited to the following:

ICP-MS, ICP-OES, FAAS, GFAA, XRF, and DCP

For the validation of analytical methods

For the preparation of "working reference samples"

For interference studies and the determination of correction coefficients

For detection limit and linearity studies

For additional intended uses, contact Technical Staff

7.0 INSTRUCTIONS FOR THE CORRECT USE OF THIS REFERENCE MATERIAL

Storage & Handling - Keep tightly sealed when not in use. Store and use at 20 ± 4°C. Do not pipet from container. Do not return portions removed for pipetting to container.

Atomic Weight; Valence; Coordination Number; Chemical Form in Solution - 51,9961; +3; 6; Cr2O72-

Chemical Compatibility - Stable in HCl, HN03, H2SO4, HF, H3PO4. Avoid basic media. Stable with most metals (Except Ba and Pb) and inorganic anions in acidic media.

Stability - 2-100 ppb levels stable for months in 1% HNO3 / LDPE container as *Total* Cr, however, the stability of this oxidation state is unknown. 1-10,000 ppm solutions chemically stable for years in 0 - 1% HNO3 / LDPE container.

Cr Containing Samples (Preparation and Solution) - Metal (soluble in HCl.); Oxides/Ores (Chrome ore/oxides are very difficult to dissolve. The following procedures [A-D] are commonly used:

- A. Fusion with KHSO4 and extraction with hot KCI. The residue fused with Na2CO3 and KCIO3, 3:1.
- B. Fusion with NaKSO4 and NaF, 2:1.
- C. Fusion with magnesia or lime and sodium or potassium carbonates, 4:1.
- D. Fusion with Na2O2 or NaOH and KNO3 or NaOH and Na2O2.

Nickel, iron, copper, or silver crucibles should be used for D. Platinum may be used for A, B and C);

Organic Matrices (Ash at 450EC followed by one of the fusion methods above or sulfuric/hydrogen peroxide acid digestions may be applicable to non oxide containing samples)

Atomic Spectroscopic Information (ICP-OES D.L.s are given as radial/axial view):

Technique/Line	Estimated D.L.	Order Type	Interferences (underlined indicates severe)
ICP-OES 205.552 nm	0.006 / 0.0008 µ	ıg/mL 1 ioı	
ICP-OES 284.325 nm	0.008 / 0.0007 µ	ıg/mL 1 ior	
ICP-OES 276.654 nm	0.01: / 0.001 µg/	mL 1 ioi	т Си, Та, <u>У</u>
ICP-MS 52 amu	40 ppt	n/a M+	36S16O, 36Ar16O - The 50Cr, 53Cr, 54Cr lines suffer
	•		from many more potential interferences from sulfur,
. *			chlorine and argon compounds of oxygen, nitrogen
•			and carbon.

- 8.0 HAZARDOUS INFORMATION Please refer to the enclosed Material Safety Data sheet for information regarding this CRM.
- 9.0 HOMOGENEITY This solution was mixed according to an in house procedure and is guaranteed to be homogeneous.

10.0 QUALITY STANDARD DOCUMENTATION





10.1 ISO 9001:2000 Quality Management System Registration - QMI Certificate Number 010105 Recognized by:

Registrar Accreditation Board (ANSI-RAB)

Standards Council of Canada (SCC)

Dutch Council for Accreditation (RVA)

Entidad Mexicana de Acreditación, a.c.(EMA)

Members of IQ Net International Certification Network:

Argentina (IRAM), Australia (QAS), Austria (ÖQS), Belgium (Avinter), Brazil (FCAV), Canada (QMI), Hong Kong (HKQAA), Columbia (ICONTEC), Czech Republic (CQS), Denmark (DS), Finland (SFS), France (AFAQ), Germany (DQS), Greece (ELOT), Hungary (MSZT), Ireland (NSAI), Israel (SII), Italy (CISQ), Japan (JQA), Korea (KSA-QA), Netherlands (KEMA); Norway (NCS), Poland(PCBC), Portugal (APCER), Singapore (PSB), Slovenia (SIQ), Spain (AENOR), Switzerland (SQS)

10.2 ISO/IEC 17025 - 1999 "General Requirements for the Competence of Testing and Calibration"

- Chemical Testing Accredited A2LA Certificate Number 883.01
- 10.3 ISO/IEC Guide 34 2000 "General Requirements for the Competence of Reference Material Producers"
 - Reference Materials Production Accredited A2LA Certificate Number 883.02

A2LA Mutual Recognition Agreement Partners:

Australia (NATA), Austria (BmwA), Belgium (BELTEST) (BKO-OBE), Canada (SCC), Chinese Taipei (CNLA), Czech Republic (NAO), Denmark (DANAK), Finland (FINAS), France (COFRAC), Germany (DAR), Hong Kong (HKAS, Ireland (NAB), Italy (SIT) (SINAL), Japan (JAB) (JNLA), Republic of Korea (KOLAS), The Netherlands (RvA), New Zealand (IANZ), Norway (NA), Portugal (IPQ), Singapore (SAC-SINGLAS), Spain (ENAC), Sweden (SWEDAC), Switzerland (SAS), United Kingdom (UKAS) and United States (NVLAP) (ICBO ES)

- 10.4 10CFR50 Appendix B Nuclear Regulatory Commission
 - Domestic Licensing of Production and Utilization Facilities
- 10.5 10CFR21 Nuclear Regulatory Commission Reporting Defects and Non-Compliance
- 10.6 MIL-STD-45662A (Obsolete/Observed)

11.0 DATE OF CERTIFICATION AND PERIOD OF VALIDITY

- 11.1 Shelf Life The period of time during which the concentration of the analyte(s) in a properly packaged, unopened, and unused standard stored under environmentally controlled and monitored conditions will remain within the specified uncertainty range. Shelf life is limited primarily by transpiration (loss of water from the solution) and infrequently, by chemical instability. Transpiration studies of chemically-stable solutions performed at the manufacturer's facility show a CRM shelf-life of twenty one months for solutions packaged in 125-mL low density polyethylene bottles. When stored under special conditions that minimize transpiration and instability, the shelf life can be extended past this limit.
- 11.2 Expiration Date The date after which a CRM should not be used. Routine laboratory use of a CRM increases transpiration losses and the chance of contamination which affect the integrity of the CRM and limit its useful life. Manufacturer concurs with state and federal regulatory agencies' recommendations that solution standards be assigned a one-year expiration date.

Certification Date:

December 06, 2006

Expiration Date:

NAMES AND SIGNATURES OF CERTIFYING OFFICERS

Certifying Officer: Paul Gaines, PhD., Senior Technical Director

Certificate Prepared By:

Nick Maida, Product
Documentation Administrator

Katalin Le, QC Manager

Certificate Approved By:

Paux Hains