



CanAg SCC EIA

**For Research Use Only.
Not for use in
diagnostic procedures.**

REF 800-85

Instructions for use. 2022-06

Read highlighted changes

EN	EXPLANATION OF SYMBOLS
BG	ОБЯСНЕНИЕ НА СИМВОЛИТЕ
CS	VÝZNAM SYMBOLŮ
DA	SYMBOLFORKLARING
DE	ERKLÄRUNG DER SYMBOLE
EL	ΕΠΕΞΗΓΗΣΗ ΤΩΝ ΣΥΜΒΟΛΩΝ
ES	SIGNIFICADO DE LOS SÍMBOLOS
ET	SÜMBOLITE SELGITUS
FR	EXPLICATION DES SYMBOLES
HR	OBJAŠNENJE SIMBOLA
HU	JELMAGYARÁZAT
IT	SPIEGAZIONE DEI SIMBOLI
LT	SIMBOLIŲ PAAIŠKINIMAI
LV	SIMBOLU SKAIDROJUMS
NL	VERKLARING DER SYMBOLEN
NO	SYMBOLFORKLARING
PL	OBJAŚNIENIE SYMBOLI
PT	EXPlicaçãO DOS SÍMBOLOS
RO	SEMNIFFICAȚIA SIMBOLURILOR
RU	ОБОНАЧЕНИЯ
SV	SYMBOLFÖRKLARING
SK	VÝZNAM SYMBOLOV
SL	RAZLAGA SIMBOLOV
SR	OBJAŠNENJE SIMBOLA
TR	SEMBOLLERİN AÇIKLAMALARI



Use By/Годно до/Použitelní do/
Holdbar til/Vervendbar bis/
Ημερομηνία λήξης/Fecha
de caducidad/Kölblik kuni/
Utiliser jusque/Rok valjanosti/
Felhasználható/Utilizzare entro/
Sunaudot iki/Izlietot idz/Houdbaar
tot/Brukes innen/Užycí przed/
Prazo de validade/Expiră la/
Использовать до/Använd före/
Použíte'né do/Uporabno do/
Upotrebljivo do/Son Kullanma Tarihi

LOT

Batch code/Номер на партида/
Číslo šarže/Lotnummer/
Chargenbezeichnung/Aριθμός
Παρτίδας/Código de lote/Partii
kood/Code du lot/Kod serije/
Sarzszám/Codice del lotto/
Partijos kodas/Partijas kods/Lot
nummer/Partikode/Kod partii/
Código do lote/Număr de lot/
Номер лота/Lotnummer/Číslo
šarže/Številka serije/Kod partie/
Parti Kodu



Date of manufacture/Дата на производство/Datum výroby/
Produktionsdato/Herstellungsdatum/
Нијеропунја парогуџић/Fecha de fabricación/Valmismatise kuupäev/
Date of fabrication/Datum proizvodnje/
Gyártási idő/Data di produzione/
Pagaminiški data/Ražošanás datums/
Productiedatum/Fremstillingstdato/
Data produkcji/Data de fabrico/Data fabrikacija/Data производства/
Tillverkningsdatum/Dátum výroby/Datum izdelave/Datum proizvodnje/Üretim tarhi

REF

Catalogue number/Katalожен номер/
Katalogove číslo/Katalognummer/
Bestellnummer/Apriθtis katalóðou/
Número de catálogo/Kataloðski broj/
Katalogússzám/Número de catalogo/
Kataloð numeris/Numur katalogð/
Catalogusnummer/Katalognummer/
Numer katalogowy/Número do catálogo/
Număr de catalog/Homer po katalogu/
Produktnummer/Katalógoð číslo/
Kataloðka številka/Kataloðski broj/
Katalog numarası



Temperature limitation/
Температурни граници/
Teplotní omezení/
Temperaturbegrenzung/
Temperaturbegrenzung/
Περιορισμό θερμοκρασίας/
Límites de temperatura/
Temperaturi piirang/
Limite de température/
Temperaturno ograničenje/
Hőmérsékletek vonatkozó korlátozás/
Limiti di temperatura/
Temperatūrini aprībojimai/
Temperatūras ierobežojums/
Temperaturbeprekning/
Temperaturbegrensninger/
Temperatury graniczne/
Limite de temperatura/
Limite de temperatūra/
Temperaturný režim/
Temperaturbegrenzung/
Teplotné obmedzenie/
Omejitev temperature/
Temperaturno ograničenje/
Sıcaklık sınırlaması/



Contains sufficient for <96> tests/Съдържа достатъчно количество за тестове
<96>Lez použit pro <96> testů/Indeholder tilstrækkeligt/Inhalt ausreichend für <96>
Prüfungen/Приєхмено етапрэс για <96> εξιτάσιc/Contenido suficiente para <96> ensayos/Kogusest piisab <96> testi läbiviimiseks/Contenu suffisante pour <96> tests/Sadrži dovoljno za <96> testova/A doboz tartalma <96> vizsgálat elvégzéséhez elegendő/Contenido suficiente per <96> saggi/Turings skirtas atlikti <96> tyrimus/Saturs pieleikams <96> testiem/Inhoud voldoende voor <96> testen/til <96> test/Tilstrekkelig inhold for <96> prøver/Wystarczy na wykonanie <96> testów/Conteúdo suficiente para <96> ensaios/Continut suficient pentru 96 de teste/Содержит достаточные количества для <96> определений/Innehåller tillräckligt till <96> antal tester/Obsah postaðaþje na tento počet testov:<96> vsebinad zadostuje za <96> testov/Sadržina dovoljna za <96> testova/<96> testleri için yeteterlik içerir



Consult Instructions for Use/
Прочетете инструкцията за употреба/Konzultujte s návodom k použití/Se brugsanvisning/Siehe Gebrauchsanweisung/Συμβούλευτείτε τις Οδηγίες σχετικά με τη χρήση/Consulte las instrucciones de uso/Vt kasutusjuhendit/Consulter le mode d'emploi/Pročítajte upute za uporabu/Olvassa el a használati utasítást/Consultare le istruzione per l'uso/Dél naudojimo Žiūrėkite instrukcijas/Izlasiet lietošanas instrukciju/Raadpleeg de instructies voor gebruik/Les instrukcene for bruk/Sprawdzić w instrukcji użycia/Consulte as Instruções de Utilização/Consultati instruções de utilizare/Обратитесь к инструкции по применению/Se bruksanvisning/Prečitate si návod na používanie/Pročítajte uputstvo za upotrebu/Kullanım Talimatlarına Bakınız

CONT

Contents of kit/Съдържание на набора/Obsah soupravy/Kittets indhold/Inhalt des Kits/Περιεχόμενα του κιτ/Contenido del kit/Komplekt sisaldað/Contenu du kit/Sadržaj opreme/A készlet tartalma/Contenuto del kit/Rinkinio turinys/Komplekta satus/Inhoud van de set/Settets innhold/Zawartość zestawu/Conteúdo do kit/Conținutul setului/Компоненты набора/Kit innehåll/Obsah úspravy/Vsebina kompleta/Sadržaj opreme/Kitin içindeler



Biological risks/Биологически опасности/Biologická rizika/Biologisk fare/Biologische Gefahren/Biولوگیک کیلðونوا/Riesgos biológicos/BioLOGIcal ohud/Risques biologiques/Biološki rizici/Biologíai kockázatok/Rischi biologici/Biologinis pavojus/Biologíkais risks/Biologische risico's/Biologiske risikoer/Zagrożenie biologiczne/Riscos biológicos/ Biologisk risk/Pericole biologice/Биологическая опасность/Biologicky rizikové/Biologické rizíká/Biolaški rizici/Biyolojik riskler

ORIG HUM

Human/C човешки производ/Лidské/Humanit/人类/διέγεντα αναφορά/ Humano/İn männitöltö/Humaine/Ljudskog porjekla/Human/Origine Umana/Žmogaus kilmës/Cilvëku izcelesmes/Human/Menneske/Ludzka/Humano/Origine umana/Человеческого происхождения/Human/L'udske/ Humanega izvora/Ljudskog porekla/İnsan

ORIG MOU

From mouse/С миши производ/Myši/Fra mus/Maus/από ποντίκι/de ratón/Hiirtel/De souris/Mišijeg porjekla/Egérből/Murino/Pelēs kilmës/No peles/Van muizen/Fra mus/Mysia/Do rato/De la soareci/Мишиного происхождения/Fran mus/Myši/Mišijeg izvora/Mišijeg porekla/Fareden

ORIG BOV

Bovine/C говеджи произход/Hovézi/Bovin/Rind/από βοοειδή/Bovino/Veistelt/Bovine/Rogate stoke/Szarvasmarha/Bovino/Jauðio/No liellopa/Bovien/Bovin/Wolowy/Bovino/Origine bovină/крунного породаго скота/Frân ko/Hovädzie/Govejega izvora/Rogate krupne stoke/Bovin



Reconstitute with/Разтворяне с/ Rozédte pomocí/Recomponere med/Rekonstituieren mit/Aνασύσταση με/Reconstituir con/Lahjendamine/Reconstituer avec/Rekonstituiraje s/Feloldáshoz/Ricostituire/con/Alturti, ištrípgant su/Atışkaidit ar/Reconstitutie met/Rekonstituores med/Odtwarzyc za pomocą/Reconstitui com/A se reconstituí cu/Raстворять в/Rekonstituera med/Rozdělte pomocou/Rekonstituiranje z/s/Ponovo formiranje sa/Yeniden oluşturulur



Manufacturer/Производител/Výrobce/Producent/Hersteller/Kataloðeuðatj/ Fabricante/Tootja/Fabricante/Producðað/ Gyártó/Fabricante/Gamtijas/Ražotájs/Fabrikant/Produsent/Producent/Fabricante/Producðator/Производитель/Tilverkare/Výrobca/Izdelovalec/Proizvodað/Üretici

CanAg SCC EIA

Instructions for use

Enzyme immunometric assay kit
For 96 determinations

**For Research Use Only.
Not for use in
diagnostic procedures.**

IMPORTANT USER INFORMATION

SCC antigen is present in skin, sweat and saliva, and is easily distributed in aerosols (e.g. as a result of sneezing). In order to avoid false elevated values due to contamination, gloves should be used after opening the kit box and throughout the test procedure when handling reagent vials, microplate, pipette tips etc.

INTENDED USE

The CanAg SCC EIA kit is intended for the quantitative determination of squamous cell carcinoma (SCC) antigen in serum.

SUMMARY AND EXPLANATION OF THE ASSAY

Squamous cell carcinoma antigen (SCCag) is a group of glycoproteins with molecular weight ~45 kDa, belonging to the family of serine/cysteine -protease inhibitors (1). The protein was originally isolated by Kato and co-workers from human squamous cell carcinoma tissue and shown to consist of at least 10 subfractions differing in isoelectric point (2). More recent studies have shown that SCC antigen is composed of two distinct but highly homologous gene products, SCCA1 and SCCA2 with different inhibitor specificities (3).

PRINCIPLE OF THE TEST

The CanAg SCC EIA is a solid-phase, non-competitive immunoassay based upon the direct sandwich technique. Calibrators and samples are incubated together with biotinylated Anti-SCC monoclonal antibody and horseradish peroxidase (HRP) labelled Anti-SCC monoclonal antibody in Streptavidin coated microstrips (4). After washing, buffered Substrate/ Chromogen reagent (hydrogen peroxide

and 3, 3', 5, 5' tetra-methylbenzidine) is added to each well and the enzyme reaction is allowed to proceed. During the enzyme reaction a blue colour will develop if antigen is present. The intensity of the colour is proportional to the amount of SCC present in the samples.

The colour intensity is determined in a microplate spectrophotometer at 620 nm (or optionally at 405 nm after addition of Stop Solution). Calibration curves are constructed for each assay by plotting absorbance value versus the concentration for each calibrator. The SCC concentrations of samples are then read from the calibration curve.

REAGENTS

- Each CanAg SCC EIA kit contains reagents for 96 tests.
- The expiry date of the kit is stated on the label on the outside of the kit box.
- Do not use the kit beyond the expiry date.
- Do not mix reagents from different kit lots.
- Store the kit at 2–8°C. Do not freeze.
- Opened reagents are stable according to the table below provided they are not contaminated, stored in resealed original containers and handled as prescribed. Return to 2–8°C immediately after use.

Component	Quantity	Storage and stability after first opening	
MICROPLA			
Microplate	1 Plate	2–8°C until expiry date stated on the plate	
	12 x 8 wells coated with Streptavidin. After opening, immediately return unused strips to the aluminium pouch, containing desiccant. Reseal carefully to keep dry.		
SCC Calibrators	5 vials, lyophilized	4 weeks at 2–8°C 3 months at -20°C	
CAL	SCC	A	1 x 0.75 mL
CAL	SCC	B	1 x 0.75 mL
CAL	SCC	C	1 x 0.75 mL

Component	Quantity	Storage and stability after first opening
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CAL SCC D 1 x 0.75 mL

CAL SCC E 1 x 0.75 mL

The lyophilised calibrators contain human SCC in a Tris-HCl buffered salt solution containing bovine serum albumin, excipient, an inert yellow dye and 0.01% methyl-isothiazolone (MIT) as preservative. To be reconstituted with water before use. **NOTE:** The exact SCC concentration is lot specific and is indicated on the label of each vial.

BIOTIN Anti-SCC

Biotin Anti-SCC 1 x 15 mL 2–8°C until expiry date stated on the vial

Biotin Anti-SCC monoclonal antibody from mouse, approximately 1 µg/mL. Contains phosphate buffered saline (pH 7.2), bovine serum albumin, bovine immunoglobulin, blocking agents, detergent, an inert blue dye and 0.01% methyl-isothiazolone (MIT) as preservative. To be mixed with Tracer, HRP Anti-SCC before use.

CONJ Anti-SCC

Tracer, HRP Anti-SCC 1 x 0.75 mL 2–8°C until expiry date stated on the vial

Stock solution of HRP Anti-SCC monoclonal antibody from mouse, approximately 40 µg/mL. Contains preservatives. To be mixed with Biotin Anti-SCC before use.

SUBS TMB

TMB HRP-Substrate 1 x 12 mL 2–8°C until expiry date stated on the vial

Contains buffered hydrogen peroxide and 3, 3', 5, 5' tetramethyl-benzidine (TMB). Ready for use.

Component	Quantity	Storage and stability after first opening
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STOP

STOP Solution 1 x 15 mL 2–8°C until expiry date stated on the vial

Contains 0.12 M hydrochloric acid. Ready for use.

WASHBUF 25X

Wash Concentrate 1 x 50 mL 2–8°C until expiry date stated on the bottle

A Tris-HCl buffered salt solution with Tween 20. Contains Germall II as preservative. To be diluted with water 25 times before use.

Indications of instability

The TMB HRP-Substrate should be colourless or slightly bluish. A blue colour indicates that the reagent has been contaminated and should be discarded.

WARNINGS AND PRECAUTIONS

For Research Use Only. Not for use in diagnostic procedures.

- Please refer to the US Department of Health and Human Services (Bethesda, Md., US) publication No. (CDC) 88-8395 on laboratory safety or any other local or national regulation.
- Handle all serum specimens as potentially infectious.
- Follow local guidelines for disposal of all waste material.

CLP (1272/2008) HAZARD CLASSIFICATION

Information about CLP (1272/2008) HAZARD CLASSIFICATION can be found at the end of this document.

SPECIMEN COLLECTION AND HANDLING

The CanAg SCC EIA is intended for use with serum. Collect blood by venipuncture and separate the serum according to common procedures. Samples can be stored at 2–8°C for 1 day. For longer periods it is recommended to store the samples at -70°C or below. Avoid repeated freezing and thawing of the samples. Allow frozen samples to thaw slowly, preferably at 2–8°C over night and then bring the samples to room temperature before analysis.

PROCEDURE

Materials required but not supplied with the kit

1. Microplate shaker

Shaking should be medium to vigorous. Longitudinal shaking approximately 200 strokes/min, oscillations 700-1100/min.

2. Microplate wash device

Automatic plate wash capable of performing 1 and 6 washing cycles with a minimal fill volume of 350 µL/well/washcycle.

An 8-channel pipette with disposable plastic tips for delivery of 350 µL is recommended if an automatic microplate washer is not used.

3. Microplate spectrophotometer

With a wavelength of 620 nm and/or 405 nm and an absorbance range of 0 to 3.0.

4. Precision pipettes

With disposable plastic tips to deliver microlitre and millilitre volumes.

An 8-channel pipette or respenser pipette with disposable plastic tips for delivery of 100 µL is useful but not essential.

5. Distilled or deionized water

For reconstitution of SCC Calibrators and for preparation of Wash Solution.

Procedural notes

1. A thorough understanding of this package insert is necessary to ensure proper use of the CanAg SCC EIA kit. The reagents supplied with the kit are intended for use as an integral unit. Do not mix identical reagents from kits having different lot numbers. Do not use the kit reagents after the expiry date printed on the outside of the kit box.

2. Reagents should be allowed to reach room temperature (20–25°C) prior to use. The assay should only be performed at temperatures between 20–25°C to obtain accurate results. Frozen specimens should be brought to room temperature slowly and must be gently but thoroughly mixed after thawing.

3. Before starting to pipette calibrators and specimens it is advisable to mark the strips to be able to clearly identify the samples during and after the assay.

4. The requirement for efficient and thorough washing for separation of bound and unbound antigen and reagents from the solid-phase bound antibody-antigen complexes is one of the most important steps in an EIA. In order to ensure efficient washing make sure that all wells are completely filled to the top edge with wash solution during each wash cycle, that wash solution is dispensed at a good flow rate, that the aspiration of the wells between and after the wash cycles is complete and that the wells are empty. If there is liquid left, invert the plate and tap it carefully against absorbent paper.

- Automatic strip washer: Follow the manufacturer's instructions for cleaning and maintenance diligently and wash the required number of wash cycles prior to and after each incubation step. It's highly recommended to use *strip* process mode and *overflow* wash mode with a dispensing volume of 800 µL. The aspiration/wash device should not be left standing with the Wash Solution for long periods, as the needles may get clogged resulting in poor liquid delivery and aspiration.

5. The TMB HRP-Substrate is very sensitive for contamination. For optimal stability of the TMB HRP-Substrate, pour the required amount from the vial to a carefully cleaned reservoir or preferably a disposable plastic tray to avoid contamination of the reagent. Be sure to use clean disposable plastic pipette tips (or respenser pipette tip).

6. Be sure to use clean disposable plastic pipette tips and a proper pipetting technique when handling samples and reagents. Avoid carry-over by holding the pipette tip slightly above the top of the well and avoid touching the plastic strip or surface of the liquid. A proper pipetting technique is of particular importance when handling the TMB HRP-Substrate solution.

Protocol Sheet

CanAg SCC EIA REF 800-85

Mix the components directly before use. Use shaking conditions according to the Instructions.

Step	Procedure		
1. Prepare SCC Calibrators A, B, C, D, E	<input type="checkbox"/> CAL	<input type="checkbox"/> SCC	Add 0.75 mL of distilled water to each vial and mix gently. Allow to stand for at least 15 minutes. NOTE: The exact concentration of each calibrator is stated on the label. This value of the calibrators should be used for calculations.
2. Prepare Wash Solution	<input type="checkbox"/> WASHBUF	<input checked="" type="checkbox"/> 25X	Dilute 50 mL of Wash Concentrate with 1200 mL of distilled water or deionized water.
3. Prepare Antibody Solution	<input type="checkbox"/> CONJ	<input type="checkbox"/> Anti-SCC	Mix 50 µL of Tracer, HRP Anti-SCC with 1 mL of Biotin Anti-SCC per strip:
Biotin Anti-SCC	No. of Strips	HRP Anti-SCC (µL)	Biotin Anti-SCC (mL)
	1	50	1
	2	100	2
	3	150	3
	4	200	4
	5	250	5
	6	300	6
	7	350	7
	9	450	9
	11	550	11
4. Wash	<input type="checkbox"/> MICROPLA		Wash each well once with Wash Solution
5. Add calibrators and samples A, B, C, D, E	<input type="checkbox"/> CAL	<input type="checkbox"/> SCC	25 µL in each well
6. Add Antibody Solution	<input type="checkbox"/> ANTI BODY SOLUTION		100 µL in each well
7. Incubate	<input type="checkbox"/> MICROPLA	1 hour shaking at room temperature	
8. Wash	<input type="checkbox"/> MICROPLA	Wash each well six times with Wash Solution	
9. Add TMB HRP-Substrate	<input type="checkbox"/> SUBS	<input type="checkbox"/> TMB	100 µL in each well
10. Incubate	<input type="checkbox"/> MICROPLA	30 min shaking at room temperature	
11. Read absorbance	<input type="checkbox"/> MICROPLA	620 nm	
Alt.11 Add Stop Solution	<input type="checkbox"/> STOP		100 µL in each well
Alt.12 Incubate	<input type="checkbox"/> MICROPLA	1 min shaking at room temperature	
Alt.13 Read absorbance	<input type="checkbox"/> MICROPLA	Read at 405 nm within 5 min	

Preparation of reagents	Stability of prepared reagent
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SCC Calibrators

4 weeks at 2–8°C
3 months at -20°C or below

Add exactly 0.75 mL of distilled water to each vial and mix gently. Allow to stand for at least 15 minutes to reconstitute. **NOTE:** The concentration of the calibrators is stated on the labels and should be used for calculation of results.

Wash Solution	2 weeks at 2–25°C in a sealed container
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Pour the 50 mL Wash Concentrate into a clean container and dilute 25- fold by adding 1200 mL of distilled or deionized water to give a buffered Wash Solution.

Antibody Solution	3 weeks at 2–8°C
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Prepare the required quantity of Antibody Solution by mixing 50 µL of Tracer, HRP Anti-SCC with 1 mL of Biotin Anti-SCC per strip (see table below and the Protocol Sheet):

No. of Strips	Tracer, HRP Anti-SCC (µL)	Biotin Anti-SCC (mL)
1	50	1
2	100	2
3	150	3
4	200	4
5	250	5
6	300	6
7	350	7
8	400	8
9	450	9
10	500	10
11	550	11
12	600	12

Be sure to use a clean plastic or glass bottle for preparation of the Antibody Solution.

Alternative: Pour the content of the Tracer, HRP Anti-SCC into the vial of Biotin Anti-SCC and mix gently. Make sure that all of the Tracer, HRP Anti-SCC is transferred to the vial of Biotin Anti-SCC.

NOTE: The Antibody Solution is stable for 3 weeks at 2–8°C. Do not prepare more Antibody Solution than will be used within this period and make sure that it is stored properly.

Assay procedure

Perform each determination in duplicate for calibrators and samples. A calibration curve should be run with each assay. All reagents and samples must be brought to room temperature (20–25°C) before use.

1. Start to prepare SCC Calibrators, Wash Solution and Antibody Solution. It is important to use clean containers. Follow the instructions carefully.
2. Transfer the required number of microplate strips to a strip frame. (Immediately return the remaining strips to the aluminium pouch containing a desiccant and reseal carefully). Wash each strip once with the Wash Solution. Do not wash more strips than can be handled within 30 min.
3. Pipette 25 µL of the SCC Calibrators (CAL A, B, C, D, E) and unknown samples (unknowns-Unk) into the strip wells according to the following scheme:

	1	2	3	4	5	6	7 etc
A	Cal A	Cal E					
B	Cal A	Cal E					
C	Cal B	Unk1					
D	Cal B	Unk1					
E	Cal C	Unk2					
F	Cal C	Unk2					
G	Cal D	Etc.					
H	Cal D						

4. Add 100 µL of Antibody Solution to each well using a 100 µL precision pipette (or an 8-channel 100 µL precision pipette). Avoid carry-over by holding the pipette tip slightly above the top of the well and avoid touching the plastic strip or the surface of the liquid.

5. Incubate the frame containing the strips for 1 hour (\pm 5 min) at room temperature (20–25°C) with constant shaking of the plate using a microplate shaker.
6. Wash each strip 6 times, using the wash procedure described in Procedural notes item 4.
7. Add 100 μ L of TMB HRP-Substrate to each well using the same pipetting procedure as in item 4. The TMB HRP-Substrate should be added to the wells as quickly as possible and the time between the addition to the first and last well should not exceed 5 min.
8. Incubate for 30 min (\pm 5 min) at room temperature with constant shaking. Avoid direct sunlight.
9. Immediately read the absorbance at 620 nm in a microplate spectrophotometer.

Option

If the laboratory does not have access to a microplate spectrophotometer capable of reading at 620 nm, the absorbance can be determined as follows:

- Alt. 9.** Add 100 μ L of Stop Solution. Mix and read absorbance at 405 nm in a microplate spectrophotometer within 5 min after addition of Stop Solution.

Measurement range

The CanAg SCC EIA measures concentrations between 0.3 and 50 μ g/L. If SCC concentrations above the measuring range are to be expected, it is recommended to dilute samples with normal human serum prior to analysis. **NOTE:** The serum used for dilution should also be measured in order to determine the endogenous SCC concentration (see "Calculation of results").

Quality control

CanChek Tumor Marker Control Sera Levels 1 and 2 (available separately, REF 107-20) are recommended for validation of the assay series. If values outside of the specified range are obtained, a complete check of reagents and reader performance should be made and the analysis repeated.

Reference material

Since no common reference material is available for SCC antigen, CanAg SCC Calibrator values are assigned against a set of in-house reference standards.

CALCULATION OF RESULTS

If a microplate spectrophotometer reader with built-in data calculation program is used, refer to the manual for the plate reader and create a program using the concentration stated on the labels of each of the SCC Calibrators.

For automatic calculation of SCC results it is recommended to use either of the following methods:

- Cubic spline curve fit method. Calibrator 0 should be included in the curve with the value 0 μ g/L.
- Spline smoothed curve fit method. Calibrator 0 should be used as plate blank.
- Interpolation with point-to-point evaluation. Calibrator 0 should be included in the curve with the value 0 μ g/L.
- Quadratic curve fit method. Calibrator 0 should be included in the curve with the value 0 μ g/L.

Note: 4-parametric or linear regression should not be used.

For manual evaluation, a calibration curve is constructed by plotting the absorbance (A) values obtained for each SCC calibrator against the corresponding SCC concentration (in μ g/L), see figure below. The unknown SCC concentrations can then be read from the calibration curve using the mean absorbance value of each specimen.

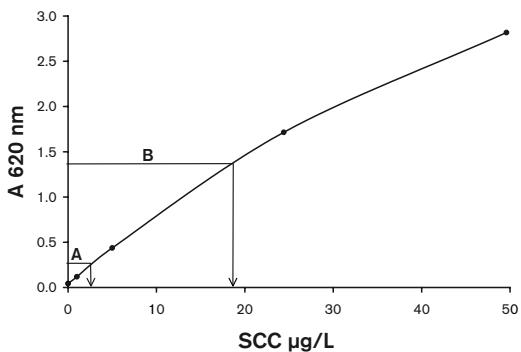
If samples in an initial analysis give SCC levels higher than 50 μ g/L the samples should be diluted 1/10 with normal human serum and reanalyzed to obtain the accurate SCC concentration. **NOTE:** The sample used for dilution should also be measured in order to determine the endogenous SCC concentration.

The SCC concentration of the undiluted sample is calculated as:

$$\text{Dilution 1/10: } 10 \times ([\text{SCC}]_{\text{Diluted sample}} - (0.9 \times [\text{SCC}]_{\text{Normal serum}}))$$

Example of results

Specimen	Calibrator values	Mean abs value (A)	SCC (µg/L)
CAL	SCC A	0 µg/L	0.043
CAL	SCC B	1 µg/L	0.119
CAL	SCC C	5 µg/L	0.437
CAL	SCC D	24 µg/L	1.715
CAL	SCC E	50 µg/L	2.818
Specimen A		0.245	2.6
Specimen B		1.363	18.3



Example (do not use this curve or table above to determine actual assay results).

LIMITATIONS OF THE PROCEDURE

SCC antigen is present in skin, sweat and saliva, and is easily distributed in aerosols (e.g. as a result of sneezing). In order to avoid false elevated values due to contamination, gloves should be used throughout the test procedure when handling reagent vials, microplate, pipette tips etc.

Anti-reagent antibodies (human anti-mouse antibody (HAMA) or heterophilic antibodies) in the sample may occasionally interfere with the assay, even though specific blocking agents are included in the buffer.

CLP (1272/2008) HAZARD CLASSIFICATION

The following warnings and precautions apply to **SUBS TMB**

Hazard pictograms:



Signal word:

Danger

Hazard Statement:

Repr. 1B: H360D May damage the unborn child.

Prevention statement:

P202 Do not handle until all safety precautions have been read and understood.

Prevention:

P280 Wear protective gloves / protective clothing / eye protection / face protection.

Precautionary statement response:

P308+P313 If exposed or concerned get medical advice/attention.

Precautionary statement disposal:

P501 Dispose of contents / container to an approved hazardous / special waste disposal facility in accordance with local and national regulations.

Restricted to professional users.

Hazardous substances: 2- Pyrrolidone

Other hazards

None of the mixtures in the kit contains any substances considered to meet the criteria classifying them as PBT and/or vPvB.

WARRANTY

The performance data presented here were obtained using the assay procedure indicated. Any change or modification of the procedure not recommended by Fujirebio Diagnostics may affect the results, in which event Fujirebio Diagnostics disclaims all warranties expressed, implied or statutory including the implied warranty of merchantability and fitness for use.

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