

Quantification of VEGF inhibitor using iLite® VEGF Assay Ready Cells

For research and professional use only. Not for use in diagnostic procedures.

This application note contains a suggested protocol and performance data. Each individual laboratory must set up their own method and perform relevant validations.

Background

Vascular Endothelial Growth Factor (VEGF) is a signalling protein which is involved in both normal vascular growth and pathological angiogenesis. Without angiogenesis, growth of solid tumors would be limited by oxygen and nutrient supply. Tumors which express VEGF can overcome this limitation and are thus able to grow and metastasize. For this reason, different anti-cancer therapies targeting VEGF have emerged, e.g. a humanized anti-VEGF antibody bevacizumab (Avastin™, Genentech) is currently widely used as a first-line therapy for colorectal cancer (1,2).

Principle of the assav

The iLite® VEGF Assay Ready Cells are engineered cells optimized to express Firefly luciferase under the control of a VEGF responsive promoter. Binding of VEGF to the VEGF receptor 2 (VEGFR2) results in activation of the VEGF regulated Firefly luciferase reporter gene construct. The Firefly luciferase signal can be measured in a luminometer following addition and incubation of luciferase substrate. The Firefly luciferase signal is proportional to the functional activity of VEGF in the sample. In the presence of inhibitory activity against VEGF, the amount of free VEGF is reduced, resulting in a decreased stimulation of Firefly luciferase production.

The Firefly luciferase signal is thus inversely proportional to the amount of inhibitory activity against VEGF in a sample. The iLite® VEGF Assay Ready Cells can therefore be utilized as a highly sensitive assay for quantification of VEGF inhibitor activity in test samples, including human serum.

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Material and equipment needed

Material and equipment	Suggested supplier	Reference
iLite® VEGF Assay Ready Cells	Svar Life Science	BM4021
Diluent (DMEM containing 9% heat inactivated FBS + 1% Penicillin-Streptomycin).	Gibco	31966-021 (DMEM) 26140-079 (FBS) 15140-122 (Penicillin- Streptomycin)
Bevacizumab	NA	NA
VEGF or analogues	Gibco	PHC9394
Firefly substrate	Promega	E2620, Bright-Glo Luciferase Assay System
Plate; White walled micro well plate suitable for luminescence	PerkinElmer	6005680
Microplate Luminometer with appropriate reading software – no filter on luminometer	Contact Svar Life Science for list of recommended suppliers	NA
Incubator, 37 °C with 5% CO ₂	NA	NA
Water bath, 37 °C	NA	NA
Single-channel and multi-channel pipettes with polypropylene disposable tips	NA	NA
Polypropylene tubes or plate for dilution	NA	NA
Single-use polypropylene reservoir	NA	NA
Plate shaker	NA	NA
Timer	NA	NA

Protocol

Preparation of VEGF inhibitor

Bevacizumab from Genentech has successfully been used to neutralize VEGF and inhibit the VEGF regulated Firefly luciferase expression in *iLite*® VEGF Assay Ready Cells (refer to the table and graph below).

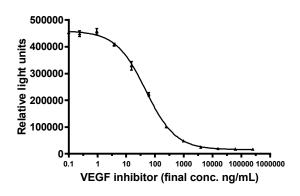


Figure 1. Example of VEGF inhibitory curve

	Bevacizumab	
Final 25 ng/mL VEGF	Suggested solution concentrations, ng/mL	
Α	1 000 000	
В	250 000	
С	62 500	
D	15 625	
E	3 906	
F	977	
G	244	
Н	61	
1	15	
J	3.8	
K	0.95	
L	0	

Table 1. Suggested calibrator solution concentrations for anti-VEGF Bevacizumab

APPLICATION NOTE



Incubation

- 1. Design a plate layout. It is recommended to perform the test at least in duplicate.
- 2. Perform a serial dilution of the reference VEGF inhibitor. Ensure matrix consistency between reference antibody solutions, control solutions, and sample solutions.
- 3. Add 20 μ L of the reference VEGF inhibitor dilutions, controls and samples to assigned wells (final concentration will be one-fourth of solution concentration).
- 4. Add 20 μL of 100 ng/ml VEGF to all wells (final concentration will be 25 ng/mL VEGF).
- 5. Place the lid on the plate, mix and incubate the plate for 30 minutes at 37 °C with 5% CO₂
- 6. Thaw the vial of *iLite*® VEGF Assay Ready Cells in a 37°C water bath with gentle agitation. The cell suspension is mixed very carefully ten times with pipette in order to ensure a homogeneous distribution of cells.
- 7. Dilute 250 µL cells with 5.75 mL Diluent.
- 8. Add 40 µL diluted cells to each well.
- 9. Place the lid on the plate, mix and incubate for 18 hours at 37 °C with 5% CO₂.

Adding substrate solutions

- 10. Equilibrate the plate and the substrate solution to room temperature.
- 11. Prepare the **Firefly luciferase** substrate according to the suppliers instructions and add 80 μL per well. Mix and protect the plate from light. After 2 minutes incubation at room temperature read in a luminometer.

Precautions

- This application note is intended for professional laboratory research use only. The data and
 results originating from following the Application Note should not be used either in diagnostic
 procedures or in human therapeutic applications.
- Use and handle the material and instruments referenced according to the supplier's/manufacturer's instructions or product specifications accompanying the individual material and instruments.
- Dispose of all sample specimens, infected or potentially infected material in accordance with good microbiological practice. All such materials should be handled and disposed as though potentially infectious.
- Residues of chemicals and preparations are generally considered as biohazardous waste and should be inactivated prior to disposal by autoclaving or using bleach. All such materials should be disposed of in accordance with established safety procedures.

Propriety Information

In accepting delivery of *iLite*® Assay Ready Cells the recipient agrees not to sub-culture these cells, attempt to sub-culture them or to give them to a third-party recipient, and only to use them directly in assays. *iLite*® cell-based products are covered by patents which are the property of Svar Life Science AB and any attempt to reproduce the delivered *iLite*® Assay Ready Cells is an infringement of these patents

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QUICK GUIDE

Quantification of VEGF inhibitor activity using iLite® VEGF Assay Ready Cells

1 Sample dilution

- Equilibrate reagents and samples to room temperature do not thaw cells and substrate reagents at this stage
- Dilute reference VEGF inhibitor, controls and samples
- Add 20 µL of ref. VEGF inhibitor solutions, controls and diluted samples to pre-assigned wells
- Add 20 µL VEGF to each well

2

• Incubate at 37 °C with 5% CO₂ for 30 min

Incubation 30 min

Add cell

- Thaw the vial of cells in a 37°C water bath. Mix cell suspension with a pipette in order to ensure a uniform cell suspension. Dilute the cells
- · Add 40 µL diluted cells to each well

4 Incubation • Incubate at 37 °C with 5% CO2 for 18 hours

ncubation 18 h

- Equilibrate the plate to room temperature
- Prepare the **Firefly luciferase** substrate according to the supplier's instructions and add 80 µL per well. Mix Protect the plate from light. After 2 min incubation read in a luminometer

5 Read plate

Troubleshooting and FAQ

Please consult the Svar Life Science website www.svarlifescience.com

References

- 1. Wang Y, Fei D, Vanderlaan M, Song A. Biological activity of bevacizumab, a humanized anti-VEGF antibody in vitro. Angiogenesis 7:335-345 (2004).
- 2. Risau W. Mechanisms of angiogenesis. Nature 386: 671-674 (1997).