

SURFE²R N1

In-depth transporter research



nanjion

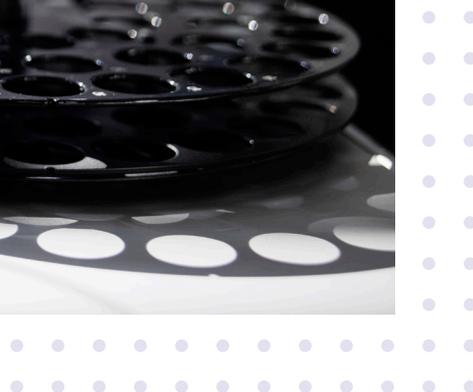
Unveiling Membrane Transport through SSM-based Electrophysiology

Nanon's SURFE²R (surface electrogenic event reader) instruments allow for functional and direct recordings of electrogenic membrane transport from symporters, exchangers, uniporters and pumps, and from ligand-gated and leak ion channels.

SURFE²R N1 facilitates in-depth functional characterization of electrogenic membrane transport in real-time with high sensitivity and high temporal resolution.

Any membrane can be accessed for recordings, ranging from isolated membrane vesicles from native tissues or cultured cells to transporters reconstituted into proteoliposomes.

It's easy to use, efficient, and unlocks completely new possibilities in transporter research and screening.



Key features at a glance

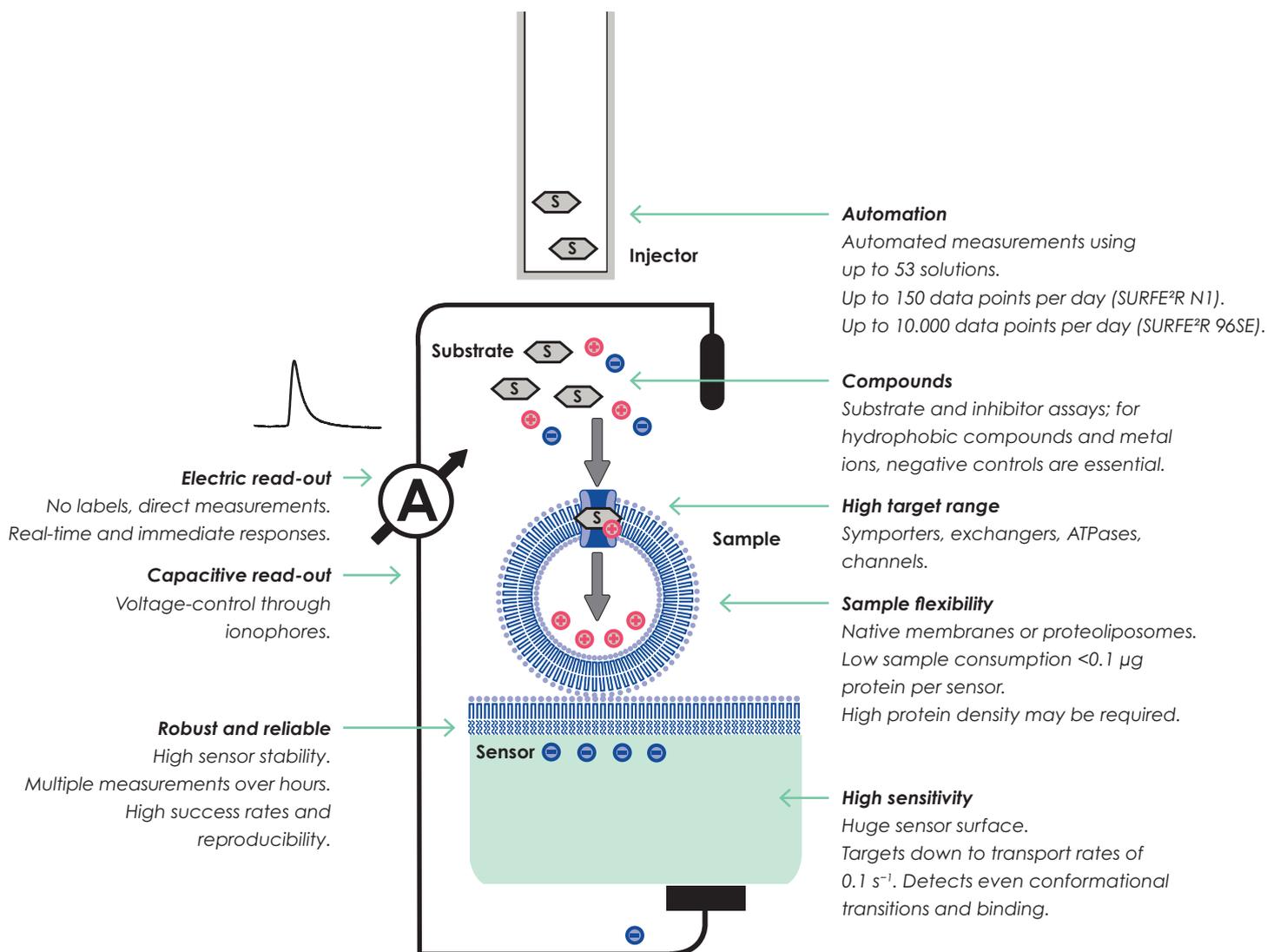
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|----------|--|----------|--|
| 1 | Functional transporter recordings in real-time | 5 | Pre-programmable, automated recordings |
| 2 | No need for labels, radioactivity or fluorescence | 6 | Broad range of validated transporter targets |
| 3 | High sensitivity & temporal resolution | 7 | Easy to learn and use — even for students |
| 4 | Access any membrane with only 0.1–1 µg protein per sensor required | 8 | Up to 150 datapoints per day |



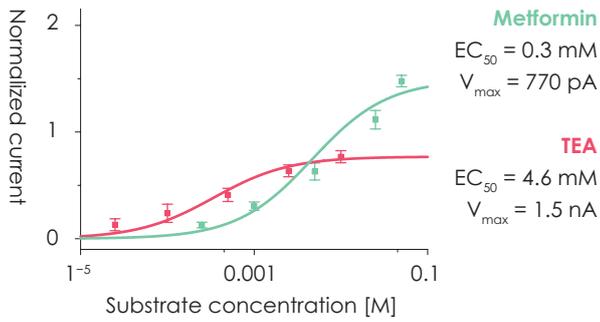
SURFE²R recording principles

Membrane fragments from cell culture, tissue or liposomes are adsorbed onto a solid supported membrane (SSM) on a gold coated sensor chip. Approximately 0.1–1 μg of membrane is required to cover the 3 mm sensor electrode, corresponding to ca 10⁹ transporter molecules. Membrane transport, i.e. charge translocation, is triggered by rapidly perfusing the sensor with a substrate or an altered ionic composition

resulting in a gradient across the membrane. The resulting electrogenic transport is measured as a transient, capacitive current which decays until a new electrochemical equilibrium has been reached. The size and shape of the current reveal information about numerous biophysical and pharmacological transporter properties like the speed of transport, coupling ratio or substrate affinity.

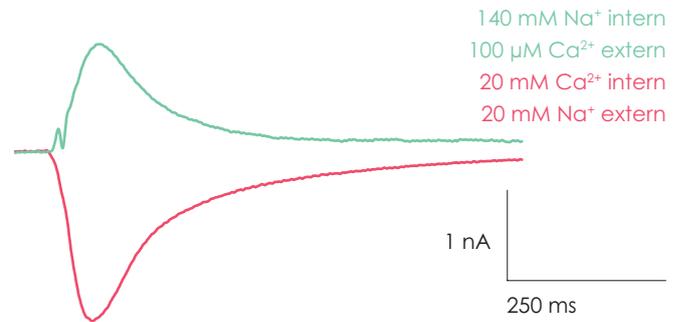


Explore the possibilities of full experimental freedom



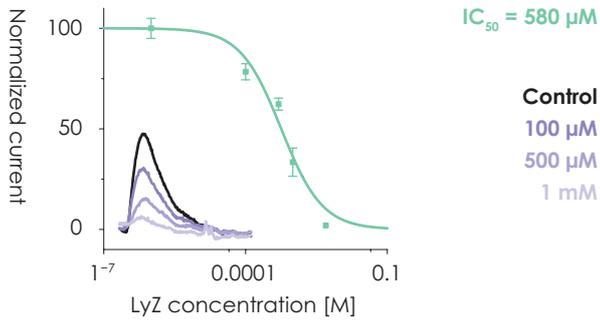
Transport assays

Substrate specificity (V_{max} , EC_{50}).



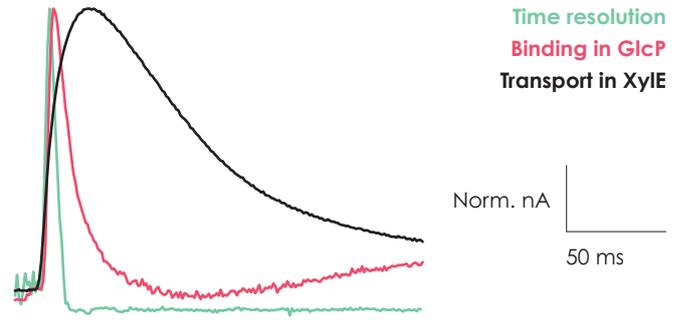
Influx and efflux assays

Symmetry of transport.



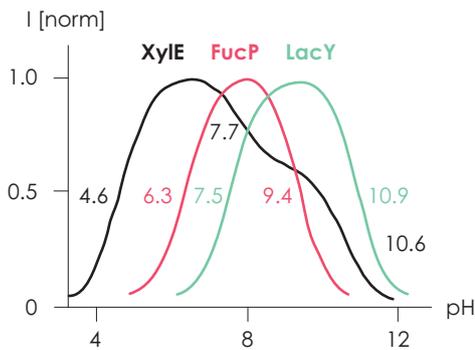
Compound assays

Inhibition and positive modulation.



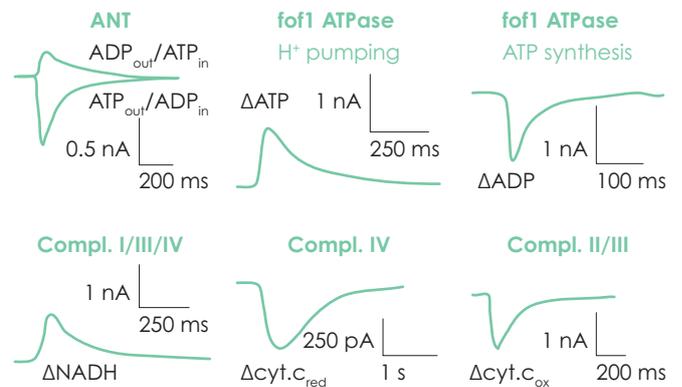
Substrate binding assay

Pre-steady state currents (k^+ , k^-).



Comparative assays

Mutants, orthologues, isoforms.



Multiple targets in one experiment

Serial recordings from the same sample and sensor.

Broad range of validated targets

Primary active transporters / Ion pumps

ATPases

NaK-ATPase
 HK-ATPase
 SERCA (Ca²⁺)
 v-ATPase (H⁺)
 FoF1-ATPase (H⁺)
 Kdp-ATPase (K⁺)
 CopA (Cu²⁺)
 ATP7A/B (Cu²⁺)
 ATP8A2 (Flippase)
 VrPPase

Redox-driven ion pumps

Complex I
 Respiratory chain complex I/III
 Respiratory chain complex II/III
 Cytochrome c-oxidase
 Respiratory chain complexes I/III/V

Light-driven ion pumps

Bacteriorhodopsin (BR)
 Rhodopsin OR1
 Rhodopsin-2 (KR2)
 Halorhodopsin (HR)
 Acerhodopsin
 Channelrhodopsin 1/2 (ChR)

Channels + Pores

Gramicidine
 P2X2
 nAChR
 A/M2
 UCP1 (Slc25a7)
 TRPC5
 TRPA1
 AQP6
 KtrAB
 TMEM175

Uniporters + Symporters + Exchangers

Inorganic ions

NhaA/B (2Na⁺/1H⁺)
 NhaP/NHA2 (1Na⁺/1H⁺)
 NCX1/mjNCX (3Na⁺/Ca²⁺)
 ecClc/Clc-3/5/7/K (Cl⁻/H⁺)
 NirC (NO₂⁻/H⁺)
 afAmt1-3/ksAmt5/ecAmtB (NH₄⁺)
 NeRh50 (NH₄⁺)
 SulP/DauA (bicarbonate)
 NIS (2Na⁺/I⁻)
 NaPi-IIb (Na⁺/PO₃²⁻)
 MntH2 (Mn(II), Zn(II), Co(II), Cd(II))
 YiiP (metals)

Sugars

SGLT1/2 (Na⁺/glucose)
 MelB (Na⁺/melibiose)
 LacY (H⁺/lactose)
 FucP (H⁺/fucose)
 XylE (H⁺/xylose)
 GlcP (glucose)
 STP10 (H⁺/glucose)

Amino acids + peptides

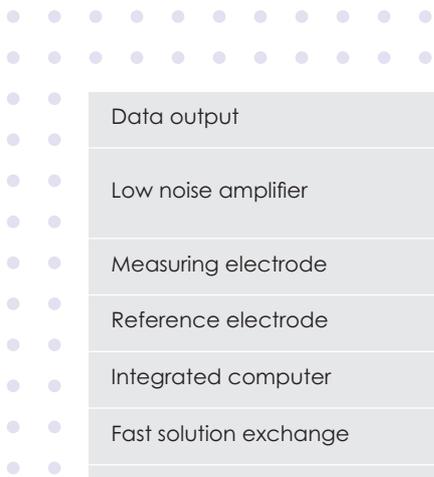
PepT1/PepT_St (di-/tri-peptides)
 YdgR/DtpA (tri-peptides)
 YhiP/DtpB (di-/tri-peptides)
 PutP (Na⁺/proline)
 GltP (Na⁺/glutamate)
 GlhTK (Na⁺/aspartate)
 EAAC1 (3Na⁺/glutamate)
 PAT1 (H⁺/proline)
 ArcD (arginine/ornithine exchange)
 CAT-IIB (cationic amino acids)
 GlyT1/2/B1 (Na⁺/glycine)
 B0AT2 (Na⁺/neutral amino acids)
 Cystin Transporter

Other organic ions

OCT1/2 (organic cations)
 OATP1B1 (organic anions)
 CNT1 (nucleosides)
 ANT (ADP/ATP exchange)
 GAT1 (Na⁺/GABA)
 BetP (betaine)
 CHT (choline)
 LicB (choline)
 NupC (H⁺/nucleoside)
 PurT (H⁺/purine)
 NaCT (Na⁺/citrate)
 SugE/Gdx (guanidinium)
 EmrE (small drugs)
 MCT13 (monocarboxylates)
 PfCRT (chloroquine/CQ)
 DgoT (H⁺/galactonate)
 Malate/lactate exchange
 Bicarbonate transport

The SURFE²R N1 includes

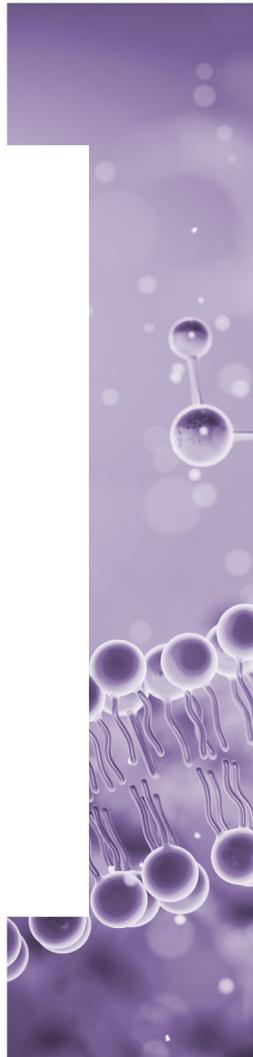
- Turn-key, stand-alone system
- PC and monitor with integrated recording and analysis software
- SURFE²R N1 Sensor 3 mm for optimal signal-to-noise
- SURFE²R N1 Sensor 1 mm for high temporal resolution
- Integrated computer with Windows and SURFE²R N1 Control software
- The SURFE²R Stimulating Optical Lid SOL (optional)
- Ph.D. level assay support and highly qualified technical support
- 1 year warranty with further optional comprehensive service plans available



Specifications

Data output	Signal amplitude: 0.1–10 nA, detection limit: 50 pA, noise: 20 pA
Low noise amplifier	Gain: 10 ⁸ –10 ¹¹ V/A, resolution: –10 V to +10 V (1.2 mV steps), RMS noise: 0.5–22 pA, low pass: 480 Hz, rise time: 0.7 ms
Measuring electrode	Re-usable gold coated sensor chip with 1 or 3 mm diameter
Reference electrode	Platinum electrode pin in measurement chamber
Integrated computer	Win11, SURFE ² R N1 Control (data recording and analysis software)
Fast solution exchange	200 µl/s, continuous flow: 0.5–6 s, time resolution: 5–30 ms
Fluidic components	2 pumps, fast switching valve, tubing cassette, IonJet with Y-fluidic
Solution handling	Autosampler with 53 positions, each ~10 ml and 2 teflon reservoirs, each ~70 ml
Solution consumption	Volume per measurement/substrate: ~300 µl (inhibitor: 1–2 ml)

accelerate your research



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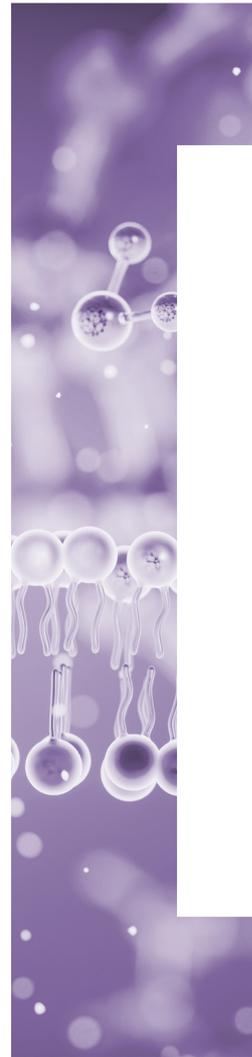
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