A synthetic mammalian electro-genetic transcription circuit
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ABSTRACT
Electric signal processing has evolved to manage rapid information transfer in neuronal networks and muscular contraction in multicellular organisms and controls the most sophisticated man-built devices. Using a synthetic biology approach to assemble electronic parts with genetic control units engineered into mammalian cells, we designed an electric power-adjustable transcription control circuit able to integrate the intensity of a direct current over time, to translate the amplitude or frequency of an alternating current into an adjustable genetic readout or to modulate the beating frequency of primary heart cells. Successful miniaturization of the electro-genetic devices may pave the way for the design of novel hybrid electro-genetic implants assembled from electronic and genetic parts.

INTRODUCTION
Recent advances in synthetic biology have led to the design of engineered cells emulating basic computational functions known from semiconductor-based electronic circuits (1–3). Synthetic gene networks implanted into living systems were able to perform tight regulation of gene expression (4), fundamental logic Boolean operations (5), establish epigenetic long-term memory (6,7), provide band-detect filter characteristics (2,8) and hysteretic signal-insulating qualities (9), program time-delayed (2) or oscillating (10) gene expression and process multichannel information within cells (11) and populations (8) as well as among different species (8,12,13). Although electronic and cell engineers are using similar circuit architectures and standardized parts (2,3) to assemble complex computing units with potentially compatible signal processing capacities, the design of electro-genetic interfaces managing mutual information processing between gene transcription in mammalian cells and electronic processing units has not yet gathered momentum. Interestingly, direct electricity-based gene expression has not been evolved as a major control theme or remains to be discovered. Previous work on electro-genetic devices mainly relied on non-specific effects of electric fields on the entire transcriptome of Escherichia coli cells (14) or on the use of electrically triggered light (15–17) to indirectly activate gene expression. In contrast to such non-specific modulation of gene expression showing genome-wide pleiotropic impact, synthetic biologists have successfully engineered mammalian transcription switches that enable reversible and adjustable activation or repression of specific transgenes in response to external stimuli, such as antibiotics (18–20), quorum-sensing molecules (21,22), (gaseous) metabolites (23–26) or cultivation temperature (27,28). All of these transcription control circuits capitalize on a generic design consisting of a synthetic transactivator (a fusion between a heterologous transcriptional repressor and a mammalian transactivation domain) which specifically binds and activates a chimeric promoter (assembled by a placing repressor-specific operator 5′ of a minimal promoter) in an inducer-responsive manner. Such a standard configuration offers optimal functional compatibility among these transcription circuits and provides a toolbox of individual transgene control modalities for assembly of complex higher order mammalian transcription networks (2,4,9,29,30).

We have designed synthetic electro-genetic devices, which enable electricity-induced expression of specific transgenes in mammalian cells as well as mammalian cell-based control of microelectronic circuits.

MATERIALS AND METHODS
Mammalian cell culture
AIRCHO-SEAP (26) transgenic for acetaldehyde-inducible expression of human placental secreted alkaline phosphatase (SEAP) was cultivated in HTS medium (Cell Culture Technology, Gravesano, Switzerland)

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supplemented with 5% (v/v) fetal calf serum (FCS, PAN Biotech GmbH, Aidenbach, Germany, cat. no. 3302, lot no. P251110), 100 U/ml penicillin, 100 μg/ml streptomycin, 1 μg/ml puromycin and 400 μg/ml G418 sulfate in a humidified atmosphere containing 5% CO₂. All expression studies were done in triplicate using 125,000 AIR/CHO-SEAP cells seeded into 2 cm² cell culture dishes. Neonatal rat ventricular cardiomyocytes were isolated from newborn rat hearts (Wistar rats; Elevage Janvier, Le Genest Saint Isle, France) as described before (31). NRCs were cultivated in 67% (v/v) Dulbecco’s Modified Eagle Medium (DMEM) (Invitrogen), 17% (v/v) M-199 EBSS (Amimed, Allschwil, Switzerland), 10% (v/v) horse serum (Invitrogen, cat. no. 16050-122, lot. 336379), 5% (v/v) FCS and 50 μg/ml gentamycin (Sigma, St Louis, MO, USA, cat. no. G1914).

**Lentiviral vectors**

Lentivectors pCD20 and pCD22, enabling acetaldehyde-responsive expression of human bone morphogenetic protein 2 (BMP-2), have been described previously (32). In brief, pCD20 (5´LTR-orisV40-cPPT-RRE-PAIR-bmp-2-3´LTR ΔU3) encodes the bmp-2 gene under control of the acetaldehyde-responsive promoter PAIR and pCD22 (5´LTR-orisV40-cPPT-RRE-PheEF1αalcR-3´LTR ΔU3) encodes constitutive expression of the acetaldehyde-dependent transactivator AlcR (26). pBP253 (5´LTR-orisV40-cPPT-RRE-PheEF1αalcR-3´LTR ΔU3) is the control lentivector for constitutive bmp-2 expression (32). Vesicular stomatitis virus-pseudotyped lentiviral particles were produced and titrated as described before (33). Abbreviations: cPPT, central polypurine tract; LTR, long-terminal repeat; orisV40, simian virus 40-derived origin of replication; PheEF1α, human elongation factor 1α promoter; RRE, rev-response element.

**Standard input device**

A custom-designed electrolysis chamber (Febikon Labortechnik, Wermelskirchen, Germany, cat. no. E-pb60) containing 250 ml 150 mM succinic acid supplemented with 1% (v/v) ethanol was used as standard input device. The chamber was placed in an airtight 3.61 polypropylene box (Migros, Zurich, Switzerland) containing a humidified atmosphere with 5% CO₂. In a typical experimental set-up, a direct current (DC) source (PowerPac HC, Bio-Rad, Hercules, CA, USA) or an alternating current (AC) generator (Lapp & Co., Zurich, Switzerland, model Th) were used to power the standard input interface for 1 h, which was then incubated for 23 h before SEAP expression was profiled using the output interface. Acetaldehyde and ethanol were quantified using Gastec tubes (26).

At high currents Joule heating \( I^2R \) of the cellular processing unit (CPU) was observed, corresponding to a temperature increase of 49 K at 500 mA (28.5 V) assuming a specific heat capacity of \( c = 4.18 \text{ kJ kg}^{-1} \text{ K}^{-1} \) for the buffer. Acetaldehyde concentrations up to 150 p.p.m. do not compromise cell viability and SEAP production (26).

**Miniaturized input device**

Modified PVDF-based hollow fibers (1 mm inner diameter, 25 mm long; Spectrum Laboratories, Rancho Dominguez, CA, USA, cat. no. M138615) were filled with 16.7 μl of a 1.7% (w/v) agarose-based hydrogel containing 125 mM succinic acid and 17% (v/v) ethanol. Both ends were equipped with platinum electrodes (diameter 1 mm), sealed and connected to a DC power source.

**Output device**

SEAP activity was converted into an electric signal using a coupled enzymatic–optical device. SEAP-containing cell culture supernatant was incubated for 5 min with p-nitrophenylphosphate as described previously (34) and the resulting p-nitrophenolate was quantitated optically at 405 nm using a Novaspec II photometer (Pharmacia, Freiburg, Germany) equipped with an output port providing an electric signal (millivolts) proportional to absorbance.

**Complementary metal-oxide semiconductor high-density microelectrode arrays**

High-density microelectrode arrays (HD-MEAs) consist of 11,016 metal electrodes and 126 channels, each of which contains recording and stimulation electronics for bidirectional communication with electrogenic cells. The HD-MEAs were manufactured as described previously (32,35). The arrays were coated with 20 μg/ml laminin for 3 h prior to seeding 105 neonatal rat cardiomyocytes (NRCs) in 1 ml medium. After incubation for 24 h, the cells were transduced with 106 pCD20/pCD22- or pBP253-derived lentiviral particles engineered for acetaldehyde-inducible or constitutive BMP-2 expression, respectively. After 24 h of transduction, the medium was exchanged and the cell-containing HD-MEAs were connected to the input interface which was set to different power levels for 1 h. The cell-containing HD-MEAs were then incubated for 48 h at 37°C before NRC beating was recorded for 20 s. The error bars represent the standard deviation from three readings.

**RESULTS AND DISCUSSION**

The generic electro-genetic input device enabling electronic transcription control in engineered mammalian cells was designed by linking electrochemical oxidation of ethanol to acetaldehyde with acetaldehyde-inducible transgene expression (26). Electrochemical oxidation of ethanol was performed using a platinum anode and cathode in an electrolysis chamber containing 1% (v/v) ethanol. Acidic pH was chosen to favour higher acetaldehyde production compared to basic pH at which the major oxidation end product is acetic acid (36). For acidification of the electrolyte, we replaced the standard perchloric acid (37) that is electrolysed to toxic chlorine with non-volatile and biocompatible succinic acid. As central cellular processing unit (CPU), we used Chinese hamster ovary (CHO-K1)-derived cells engineered for constitutive expression of the *Aspergillus nidulans*-derived acetaldehyde-dependent...
transactivator AlcR which binds and activates the acetaldehyde-inducible promoter \( \text{P}_{\text{AIR}} \) in the presence of acetaldehyde and so triggers expression of the human placental SEAP (26). SEAP production was scored by an enzymatic–optical process, consisting of a photodiode converting SEAP-catalysed dephosphorylation of \( p \)-nitrophenylphosphate to coloured \( p \)-nitrophenolate into a dose-dependent electric signal thereby providing a standard gene-electronic output device (Figure 1a). Input and output devices could either be operated as standalone electronic-cell interfaces or be sequentially linked to use engineered mammalian cells as CPUs plugged into an integrated electronic circuit.

When input devices and CPUs were operated with DC < 30 mA no significant SEAP production could be observed. Between 30 mA and 140 mA there was a direct correlation between input current, acetaldehyde production and transgene expression (Figure 1b and c). Between 140 mA and 200 mA DC SEAP expression reached a plateau since acetaldehyde-inducible transgene expression became saturated although the input device continued to produce increasing acetaldehyde levels. Beyond 200 mA DC SEAP production decreased as a consequence of a current-induced temperature increase, which steadily reduced CPU viability (Figure 1b; Materials and Methods section). Such thermal destruction is a characteristic the electro-genetic device shares with any electronic circuitry.

The photodiode-based output device was shown to provide electric signals, which were proportional to SEAP production in a range of 0–30 U/L (corresponding to an output signal of 0–1000 mV), which enabled full coverage of the transgene expression dynamics of mammalian cells (Figure 1b and d). Consolidating the dose-response characteristics of input and output devices as well as the CPU, the entire synthetic electro-genetic circuitry shows an overall dynamic range of 30–140 mA for the input current and 0.7–10 mV for the output signal.

A mammalian cell-based integrator

Within its linear operation range (30 mA DC ≤ input current ≤ 140 mA DC, see above), the electro-genetic device functions as a cell-based electronic integrator mimetic since electrochemical acetaldehyde production is a direct function of the exposure time and the intensity of the current. For validation of the integrator characteristics, the electro-genetic device was connected for different periods of time to currents with varying intensities, while the overall amount of electron flux was kept constant (\( I \times t = \) constant). The observation that acetaldehyde levels as well as the electric output were identical for all time profiles, suggests that the device has the capacity to process electronic signals in an integrator-like manner (Figure 2a).

A mammalian cell-based amplitude-modulation (AM) detector

The CPU’s electronic signal integration capacity is not limited to DC but can also be operated to score the amplitude of an AC power source. Upon connection of the CPU to an AC of 50 Hz, the resulting acetaldehyde

\[
\text{SEAP Production (x 0.25 U/L)}
\]

\[
\text{Input Signal (mA)}
\]

\[
\text{Output Signal (mV)}
\]

\[
\text{Voltage (V)}
\]

\[
\text{Current Intensity (mA)}
\]

\[
\text{Output Signal (ipm)}
\]

\[
\text{SEAP Activity (U/L)}
\]

Figure 1. Synthetic mammalian cell-based electro-genetic device. (a) Circuit diagram of the electro-genetic device. DC applied to the input device results in the electrochemical conversion of ethanol into acetaldehyde, which enables the acetaldehyde-dependent transactivator AlcR to bind and induce transcription from its cognate promoter \( \text{P}_{\text{AIR}} \) which triggers transcription of the human placental SEAP. SEAP subsequently catalyses the production of coloured \( p \)-nitrophenolate, which is quantified photometrically at 405 nm by a photodiode and converted into an electric output signal. AlcR, acetaldehyde-inducible transactivator; \( \text{P}_{\text{AIR}} \), polyadenylation signal; \( \text{P}_{\text{AIR}} \), AlcR-responsive promoter; \( \text{P}_{\text{LTER}} \), murine stem cell virus 5′ long terminal repeat-derived promoter. (b) Characterization of the CPU. The CPU was connected to DC of different intensities for 1 h, and the resulting acetaldehyde concentration as well as the electric output signal were quantified after 24 h. (c) Correlation between DC input and corresponding voltage. (d) Characterization of the output interface. SEAP activity was plotted against the electric signal measured by the output interface.
production as well as the electric signal generated by the output interface were monitored and were shown to directly correlate with the amplitude of the electric input signal thereby confirming the ability of the electro-genetic device to function as an AM detector (Figure 2).

A cell-based frequency modulation detector

The efficacy of AC-based electrolysis is frequency dependent—with increasing frequencies, polarization phenomena at the electrodes occur and re-oxidation/re-reduction of the reaction products further decrease the electrochemical production of acetaldehyde in a frequency-dependent manner (Figure 2c). In order to characterize the ability of the electro-genetic device to detect the frequency of an AC and to convert it into a DC output signal, the input interface of the electro-genetic device was connected to AC with constant amplitude (50 mA) and increasing frequencies (50–10 000 Hz). With increasing AC frequencies, decreasing acetaldehyde production was dose-dependently translated into decreasing SEAP expression, which converted into a linear electric signal by the enzymatic–optical output interface thereby

![Figure 2](image-url)

**Figure 2.** Electro-genetic circuits. (a) Mammalian cell-based integrator. The CPU was connected to DC of different intensities and for various periods of time; the product of time and current intensity was kept constant (t × I = constant). The resulting acetaldehyde levels and electric output signals were scored after 24 h. (b) Mammalian cell-based AM detector. The CPU was connected to an AC of 50 Hz and different intensities for 1 h and the acetaldehyde levels as well as the electric output current were quantified after 24 h. (c) Mammalian cell-based FM detector. The CPU was connected to an AC of 50 mA and different frequencies for 1 h and the acetaldehyde levels as well as the electric output current were quantified after 48 h.

![Figure 3](image-url)

**Figure 3.** Mammalian cell-based frequency generator. (a) Circuit diagram of the cell-based frequency generator. DC power converts ethanol into acetaldehyde, which dose-dependently triggers expression of the BMP-2 in engineered rat cardiomyocytes (AIRNR-C-BMP-2) and increases the contraction frequency (tachycardia). (b) The beating frequency of cardiomyocytes is recorded as a function of the input current and acetaldehyde concentration using a CMOS-based HD-MEA. NC: negative control, mock-transduced cardiomyocytes; PC: positive control, cells transduced for constitutive BMP-2 expression. (c) HD-MEA-based analysis of the electrogenic behaviour of NRCs engineered for electro-inducible acetaldehyde-responsive BMP-2 expression. The dataset shown as example was recorded at a direct input current of 50 mA corresponding to a beating frequency of 2.1 Hz. Detailed activation map illustrating the average signal shape of the 121 selected electrodes during 10 s. Average signal shape over all 121 channels. Raster plot showing a dot for each contraction on each channel over time. Inter-burst interval or inter-beat interval used to calculate the average beating frequency and beating frequency variation. (d) Zoom-in of two selected bursts/beats on different electrodes. (e) Long signal trace showing the synchronized contraction frequency on five selected electrodes or channels.
validating the electro-genetic device as a frequency modulation (FM) detector (Figure 2c).

**A cell-based frequency generator**

Electric signals linked to complex intracellular signalling cascades are well-known to manage muscular contraction in specialized mammalian cells (38). For example, cardiac ventricular contraction frequency of NRCs is modulated by BMP-2 that induces receptor-mediated activation of the myocyte-specific enhancer factor 2A via phosphatidylinositol 3-kinase in a dose-dependent manner (39). By transducing NRCs cultivated on complementary metal-oxide semiconductor (CMOS)-based HD-MEAs with lentiviral particles (32) engineered for electro-inducible acetaldehyde-responsive expression of BMP-2, we were able to convert DC into an oscillating electronic signal with a defined frequency (Figure 3a). Challenging this cell-based frequency generator with increasing input current resulted in elevated BMP-2 expression which stimulated NRCs to beat at higher frequency as scored by the HD-MEA (Figure 3b–e).

**Miniaturization of the input interface**

Akin to electronic equipment, which is continuously miniaturized, we have reduced the size of the initial electro-genetic input device by a factor of 15000 (electrolyte volume: 16.7 µl versus 250 ml). To achieve this reduction factor, we have cast a 17% (v/v) ethanol-containing hydrogel into a PVDF-based hollow fiber membrane which was powered via two platinum electrodes (Figure 4a). Application of an electric input between 1 µA and 4 µA (5–15 V) produced a dose-dependent output signal when linked to a miniaturized CPU (mCPU) (Figure 4b). As with electronic devices, miniaturization of the electro-genetic control device reduced power consumption to achieve maximum transgene expression levels by 30,000-fold (60 µW, compared to 1.8 W of a standard CPU).

Although microelectronic devices are dominating our daily life and control most of the analytical instruments advancing life science research, our molecular understanding of how electricity impacts biological function remains largely limited to specialized electrogenic cells such as nerve and muscle cells and appropriate applications such

![Figure 3. Continued.](http://nar.oxfordjournals.org/)
as the pacemaker. The electro-genetic device described here provides a first example of modulating transgene expression in response to an electric current by coupling an electricity-triggered electrochemical reaction to a synthetic gene network engineered into mammalian cells. Such electro-genetic devices may define novel interfaces between microelectronic and biological transcription circuits and such electro-genetic information crosstalk may one day control therapeutic transgene expression or process disease signals of prosthetic implant devices thereby harnessing the full potential of progress in electronics sectors for human therapy.

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