# Package 'BrainMetabolism'

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Title calculate brain metabolism rates from extracellular conentrations	
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<b>Depends</b> R (>= 3.1.0), ggplot2, stats	
Imports ggplot2	
<b>Description</b> calculate CMRO2, CMRgluc, mitochondrialPO2 provide functions to process data from biosensors	
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BrainMetabolism-package

calculate brain metabolism rate from extracellular conentrations

#### **Description**

calculate CMRO2, CMRgluc, mitochondrialPO2 ... provide functions to process data from biosensors

#### **Details**

Package: BrainMetabolism

Type: Package
Version: 1.0
Date: 2015-03-03
License: MIT

#### Author(s)

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## References

Piilgaard Lauritzen JCBFM (2009) 29, 1517 Gjedde et al JCBFM (2005) 25(9), 1183 Gjedde et al JCBFM (2000) 20(4), 747 Du et al JCBFM 2012 32(9) Gruetter et al J Neurochem 1998 70(1) balanca et al (2016) in preparation

## **Examples**

```
MyTP02<-23 #mmHg
MyLDF<-95 #percent
MyL<-L.calc(CMR=219, TP02=MyTP02, P50=36, h=2.7, Ca=8, cbf=53 )
MyCMR02<-CMR02.calc(LDF=MyLDF, MyTP02, P50=36, h=2.7, Ca=8, L=MyL,cbfbase=53)

MyGlucBrain<-1 #mM
MyGlucPlasma<-6 #mM
MyCRMgluc<-CMRGluc.calc(MyGlucbrain, Vd=0.77, Kt=13.4, Tmax=1.35, Gplasma=MyGlucPlasma)</pre>
```

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auc

area under the curve

#### **Description**

take a numeric vector and return the area under the curve (AUC)

## Usage

```
auc(data, pos = TRUE)
```

## Arguments

data : a numeric vector

pos : a logical value. if TRUE (default) AUC is calculated on positive values, if

FALSE on negative values.

#### Value

auc: area under the curve

max/min: maxiaml value (or minimal if pos=FALSE)

calibration

Sensor calibration

## **Description**

Fit an  $n^{th}$  degree polynom

$$(y = C_n X^n + C_{n-1} X^{n-1} + \ldots + C_1 X + C_0)$$

to vectors x=volt and y=concentration

## Usage

```
calibration(volt, mol, order = 1)
```

## **Arguments**

volt : a numeric vector of the biosensor voltage

mol : a numeric vector of the molecule conentration in the medium (x and y must

have the same length)

order : a numeric value for the polynomial degree. default is one.

#### Value

Coef: coeficients in ascending order (i.e. C0, C1, C2, ..., Cn)

R2: goodness of fit

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CMRGluc.calc

CMRGluc calculation

## Description

take extracellular glucose concentration and return brain metabolic rate of glucose, using a reversible Michaelis-Menten model equation:

## Usage

```
CMRGluc.calc(Gbrain, Vd = 0.77, Kt = 1.4, Tmax = 1.27, Gplasma = 7.3)
```

## **Arguments**

Gbrain : a vector/numeric value of extracellular glucose concentration in mmol/L, time

decay

Vd : glucose brain space diffusion (default is 0.77 ml/g)

Kt : glucose apparent maximal transport rate (default is 1.4 mmol/L)

Tmax : the apparent maximal transport rate (default is 1.27 micormol/g/min)

Gplasma : plasma glucose concentration (default=6mmol/L)

## **Details**

$$G_{brain} = V_d \frac{\left(\frac{T_{max}}{CMR_{gluc}} - 1\right) \times G_{plasma} - K_t}{\frac{T_{max}}{CMR_{gluc}} + 1}$$

#### Value

CMRGlucose: numeric value of Cerebral metabolic rate of glucose in micromol/g/min

#### References

Morgenthaler et al. Neurochem Int 2006, 48

Ori et al. Anesthesiology 1986, 65(2)

Duarte et al. Front in Neuroenrgetics 2009, 1

Gruetter et al J Neurochem 1998, 70(1)

CMRO2.calc 5

CMR02.calc	CMRO2 calculation
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#### **Description**

take tissue oxygen pressure (tPO2) and cerebral blood flow (relative value, e.g. laser doopler lowmetry BPU)

### Usage

```
CMRO2.calc(LDF, TPO2, P50 = 36, h = 2.7, Ca = 8, L = 4.03, cbfbase = 53)
```

### **Arguments**

LDF : a vector/numeric value of LDF (percentage of baseline, i.e: basal value is 100

%)

TPO2 : a vector/numeric value of brain oxygnen pressure (mmHg)
P50 : half-saturation tension of hemoglobine (default is 36 mmHg)

h : hill's coeficient

ca : oxygen arterial concentration (default is 8 micromol/ml)

: effective diffusion coefficient of oxygen in brain tissue, default is 4.03 mi-

comol/100g/mmHg, but one should use the L.calc function to calculate it from

their data.

cbfbase : basal expected value of CBF (default is 53 ml/100g/min wich was used to

calculate L)

LBF and TPO2 must be the same length

#### **Details**

$$PbtO_2 = P_{50} \cdot \sqrt[h]{\frac{2 \cdot C_a \cdot CBF}{CMRO_2} - 1} - \frac{CMRO_2}{2 \cdot L}$$

#### Value

CMRO2: vector/numeric value of Cerebral Metabolic Rate of Oxygen in micromol/100g/min

#### References

Gjedde et al JCBFM (2005) 25(9), 1183 Piilgaard et al JCBFM (2009) 29, 1517 6 correction.TPO2

correction.Temp

Temperature correction

#### Description

Biosensor enzymatic reaction, that underpin amperometric measures, has a sigmoid relation to temperature :

$$m(x, P) = \frac{P_1 + (P_2 - P_1)}{(1 + \exp((P_3 - x)/P_4))}$$

parameters are different for each enzyme and has been measured in vitro.

This function correct biosensor signal depending on temperature conditions during calibration and experimentation

## Usage

```
correction. Temp(x, enz, temp.calib = 25, temp.exp = 37)
```

#### **Arguments**

x : a numeric vector of the biosensor voltage

enz : enzyme on the biosensor i.e. "glucose", "lactate", "glutamate, "daao"

temp.calib : temperature of the medium where sensor has been calibrated the default is

25°C (i.e. room temperature).

temp.exp : temperature of the medium during experiment the default is 37°C (i.e. animal

central temperature)

#### Value

volt.temp.cor: a vector of corected x values for temperature

#### References

balanca et al 2015

correction.TP02

Oxygen Tension correction

#### **Description**

Biosensor enzymatic reaction, that underpin amperometric measures, has an asymptotic relation to oxygen tention in the medium :

$$m(PO_2, P) = P_1 + (P_2 - P_1) \times \exp(-\exp(P_3)PO_2)$$

parameters are different for each enzyme and has been measured in vitro

This function correct biosensor signal depending on PO2 conditions during calibration and experimentation

IO2.calc 7

#### Usage

```
correction.TP02(x, enz, TP02 = 28)
```

#### **Arguments**

x : a numeric vector of the biosensor voltage

enz : enzyme on the biosensor i.e. "glucose", "lactate", "glutamate, "daao"

TP02 : oxygene tension in the medium during the experiment. the default is 30mmHg

measured in anesthetized rat brain.

#### Value

volt.O2.cor: a vector of corected x values for TPO2

#### References

balanca et al 2015

IO2.calc

Oxidative index calculation

#### **Description**

take CMRO2 and LDF to give an oxidative index

#### Usage

```
IO2.calc(CMRO2, LDF, cbfbasal = 53)
```

#### Arguments

CMRO2 : vector/numeric value of Cerebral metabolic rate of oxygen in micromol/100g/min

LDF : a vector/numeric value of LDF (percentage from baseline, baseline is 100 %)

cbfbasal : basale expected value of CBF from the litterature (default is 53 ml/100g/min

wich was used to calculate L)

#### Details

$$IO2 = CMRO2/(cbfbasal * LDF)$$

#### Value

IO2: oxidative index, reflect the degree of flow metabolism coupling

## References

Gjedde et al JCBFM (2000) 20(4), 747

8 L.calc

L.calc Calculate the effective diffusion coefficient of oxygen in brain tissue, L.

## **Description**

take CMRO2 and cerebral blood flow (CBF) to calculate the effective diffusion coefficient of oxygen in brain tissue (L)

## Usage

L.calc(CMR = 219, TPO2, P50 = 36, 
$$h = 2.7$$
, Ca = 8, cbf = 53)

## **Arguments**

CMR : Cerebral Metabolic Rate of Oxygen, default is 219 micro mol/100

TPO2 : numeric value of brain oxygnen pressure (mmHg)

P50 : half-saturation tension of hemoglobine (default is 36 mmHg)

h : hill's coeficient

Ca : oxygen arterial concentration (default is 8 micromol/ml)

cbf : expected value of CBF (default is 53 ml/100g/min wich was used to calculate

Τ.

#### **Details**

$$PbtO_2 = P_{50} \cdot \sqrt[h]{\frac{2 \cdot C_a \cdot CBF}{CMRO_2} - 1} - \frac{CMRO_2}{2 \cdot L}$$

#### Value

L: numeric value of the effective diffusion coefficient of oxygen in brain tissue micomol/100g/mmHg

#### References

Gjedde et al JCBFM (2005) 25(9), 1183

Piilgaard et al JCBFM (2009) 29, 1517

noise.na 9

noise.na Remove artifacts from biosensor signal, based on data's standar deviation

## **Description**

Remove artifacts from biosensor signal, based on data's standar deviation

#### Usage

```
noise.na(data, z = 20, width = 30)
```

## Arguments

data : a numeric vector

z : a numeric value. Number of SD over which values should be exculded

width : size of the window used to roll SD over data (see roll.funct)

#### Value

a vector with NA remplacing exculded values

OGI.calc

Oxygene Glucose index (OGI)

## Description

take CMRO2 and CMRGlucose to give an OGI

## Usage

```
OGI.calc(CMRO2, CMRGluc)
```

## Arguments

cmr02 : vector (numeric value) of Cerebral metabolic rate of oxygen in micromol/100g/min

CMRGluc : a vector (numeric value) of CMRGlucose in micromol/100g/min

## **Details**

$$OGI = CMRO2/CMRGlucose$$

#### Value

**OGI** 

10 polyval

polyfit

polynomial fit

## **Description**

Fit a n<sup>th</sup> degree polynom

$$(y = C_n X^n + C_{n-1} X^{n-1} + \ldots + C_1 X + C_0)$$

to vectors x and y

## Usage

```
polyfit(x, y, order = 1)
```

## Arguments

x : a numeric vector

y : a numeric vector (x and y must have the same length) order : a numeric value for the polynomial degree. default is one.

## Value

model formula

Coef: coeficients in ascending order (i.e. C0, C1, C2, ..., Cn)

R2: Rsquare

polyval

polynomial evaluation

## **Description**

evaluation a  $n^{th}$  degree polynom

$$(y = C_n X^n + C_{n-1} X^{n-1} + \ldots + C_1 X + C_0)$$

at given values of x

## Usage

## Arguments

: a numeric vector

coef : a n dimention vector corresponding to the polynom coefficient (ascending or-

der, i.e.  $C0, C1, C2, \dots, Cn$ )

roll.funct

## Value

y

roll.funct

apply a function FUN on a rooling windows of a vector

## Description

apply a function FUN on a rooling windows of a vector

## Usage

```
roll.funct(data, width, FUN, size = T, ...)
```

## Arguments

data : a numeric vector

width : the size of the rolling window

FUN : the function to apply

size : a logical value indicating if the returned vector have the same length as original

data. default is TRUE.

... : additional argument to pass to the FUN

## Value

a vector with FUN result

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