

# Package ‘MCID’

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**Type** Package

**Title** Estimating the Minimal Clinically Important Difference

**Version** 0.1.0

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**Description** Apply the marginal classification method to achieve the purpose of providing the point and interval estimates for the minimal clinically important difference based on the classical anchor-based method. For more details of the methodology, please see Zehua Zhou, Leslie J. Bisson and Jiwei Zhao (2021) <[arXiv:2108.11589](#)>.

**License** GPL (>= 2)

**Encoding** UTF-8

**Imports** stats

**RoxygenNote** 7.1.0

**NeedsCompilation** no

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cv.imcid	<i>Selection of the tuning parameters for determining the MCID at the individual level</i>
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### Description

cv.imcid returns the optimal tuning parameter  $\delta$  and  $\lambda$  selected from a given grid by using k-fold cross-validation. The tuning parameters are selected for determining the MCID at the individual level

### Usage

```
cv.imcid(x, y, z, lamseq, delseq, k = 5, maxit = 100, tol = 0.01)
```

### Arguments

x	a continuous variable denoting the outcome change of interest
y	a binary variable denoting the patient-reported outcome derived from the anchor question
z	a vector or matrix denoting the patient's clinical profiles
lamseq	a vector containing the candidate values for the tuning parameter $\lambda$ , where $\lambda$ is the coefficient of the penalty term, used for avoiding the issue of model overfitting
delseq	a vector containing the candidate values for the tuning parameter $\delta$ , where $\delta$ is used to control the difference between the 0-1 loss and the surrogate loss. We recommend selecting the possible values from the neighborhood of the standard deviation of x
k	the number of groups into which the data should be split to select the tuning parameter $\delta$ by cross-validation. Defaults to 5
maxit	the maximum number of iterations. Defaults to 100
tol	the convergence tolerance. Defaults to 0.01

### Value

a list including the combinations of the selected tuning parameters and the value of the corresponding target function

### Examples

```
n <- 500
lambdaseq <- 10 ^ seq(-3, 3, 0.1)
deltaseq <- seq(0.1, 0.3, 0.1)
a <- 0.1
b <- 0.55
c <- -0.1
```

```

d <- 0.45

set.seed(721)
p <- 0.5
y <- 2 * rbinom(n, 1, p) - 1
z <- rnorm(n, 1, 0.1)
y_1 <- which(y == 1)
y_0 <- which(y == -1)
x <- c()
x[y_1] <- a + z[y_1] * b + rnorm(length(y_1), 0, 0.1)
x[y_0] <- c + z[y_0] * d + rnorm(length(y_0), 0, 0.1)

sel <- cv.imcid(x = x, y = y, z = z, lamseq = lambdaseq,
               delseq = deltaseq, k = 5, maxit = 100, tol = 1e-02)
sel$'Selected lambda'
sel$'Selected delta'

```

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cv.pmcid

*Selection of the tuning parameter for determining the MCID at the population level*


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### Description

cv.pmcid returns the optimal tuning parameter  $\delta$  selected from a given grid by using k-fold cross-validation. The tuning parameter is selected for determining the MCID at the population level

### Usage

```
cv.pmcid(x, y, delseq, k = 5, maxit = 100, tol = 0.01)
```

### Arguments

x	a continuous variable denoting the outcome change of interest
y	a binary variable indicating the patient-reported outcome derived from the anchor question
delseq	a vector containing the candidate values for the tuning parameter $\delta$ , where $\delta$ is used to control the difference between the 0-1 loss and the surrogate loss. We recommend selecting the possible values from the neighborhood of the standard deviation of x
k	the number of groups into which the data should be split to select the tuning parameter $\delta$ by cross-validation. Defaults to 5
maxit	the maximum number of iterations. Defaults to 100
tol	the convergence tolerance. Defaults to 0.01

**Value**

a list including the selected tuning parameter and the value of the corresponding target function

**Examples**

```
n <- 500
deltaseq <- seq(0.1, 1, 0.1)
a <- 0.2
b <- -0.1
p <- 0.5

set.seed(115)
y <- 2 * rbinom(n, 1, p) - 1
y_1 <- which(y == 1)
y_0 <- which(y == -1)
x <- c()
x[y_1] <- rnorm(length(y_1), a, 0.1)
x[y_0] <- rnorm(length(y_0), b, 0.1)

sel <- cv.pmcid(x = x, y = y, delseq = deltaseq, k = 5,
               maxit = 100, tol = 1e-02)
sel$'Selected delta'
sel$'Function value'
```

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imcid

*Point and interval estimation for the MCID at the individual level*


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**Description**

We formulate the individualized MCID as a linear function of the patients' clinical profiles. `imcid` returns the point estimate for the linear coefficients of the MCID at the individual level

**Usage**

```
imcid(x, y, z, n, lambda, delta, maxit = 100, tol = 0.01, alpha = 0.05)
```

**Arguments**

<code>x</code>	a continuous variable denoting the outcome change of interest
<code>y</code>	a binary variable indicating the patient-reported outcome derived from the anchor question
<code>z</code>	a vector or matrix denoting the patient's clinical profiles
<code>n</code>	the sample size
<code>lambda</code>	the selected tuning parameter $\lambda$ , can be returned by <code>cv.imcid</code>
<code>delta</code>	the selected tuning parameter $\delta$ , can be returned by <code>cv.imcid</code>
<code>maxit</code>	the maximum number of iterations. Defaults to 100

tol                    the convergence tolerance. Defaults to 0.01  
 alpha                 nominal level of the confidence interval. Defaults to 0.05

### Value

a list including the point estimates for the linear coefficients of the individualized MCID and their standard errors, and the corresponding confidence intervals based on the asymptotic normality

### Examples

```
n <- 500
lambdaseq <- 10 ^ seq(-3, 3, 0.1)
deltaseq <- seq(0.1, 0.3, 0.1)
a <- 0.1
b <- 0.55
c <- -0.1
d <- 0.45
### True linear coefficients of the individualized MCID: ###
### beta0=0, beta1=0.5 ###

set.seed(115)
p <- 0.5
y <- 2 * rbinom(n, 1, p) - 1
z <- rnorm(n, 1, 0.1)
y_1 <- which(y == 1)
y_0 <- which(y == -1)
x <- c()
x[y_1] <- a + z[y_1] * b + rnorm(length(y_1), 0, 0.1)
x[y_0] <- c + z[y_0] * d + rnorm(length(y_0), 0, 0.1)
sel <- cv.imcid(x = x, y = y, z = z, lamseq = lambdaseq,
               delseq = deltaseq, k = 5, maxit = 100, tol = 1e-02)
lamsel <- sel$'Selected lambda'
delsel <- sel$'Selected delta'
result <- imcid(x = x, y = y, z = z, n = n, lambda = lamsel,
               delta = delsel, maxit = 100, tol = 1e-02, alpha = 0.05)
result$'Point estimates'
result$'Standard errors'
result$'Confidence intervals'
```

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pmcid

*Point and interval estimation for the MCID at the population level*

---

### Description

pmcid returns the point estimate for the MCID at the population level

**Usage**

```
pmcid(x, y, n, delta, maxit = 100, tol = 0.01, alpha = 0.05)
```

**Arguments**

<code>x</code>	a continuous variable denoting the outcome change of interest
<code>y</code>	a binary variable indicating the patient-reported outcome derived from the anchor question
<code>n</code>	the sample size
<code>delta</code>	the selected tuning parameter $\delta$ , can be returned by <code>cv.p<code>mcid</code></code>
<code>maxit</code>	the maximum number of iterations. Defaults to 100
<code>tol</code>	the convergence tolerance. Defaults to 0.01
<code>alpha</code>	nominal level of the confidence interval. Defaults to 0.05

**Value**

a list including the point estimate of the population MCID and its standard error, and the confidence interval based on the asymptotic normality

**Examples**

```
n <- 500
deltaseq <- seq(0.1, 1, 0.1)
a <- 0.2
b <- -0.1
p <- 0.5
### True MCID is 0.5 ###

set.seed(115)
y <- 2 * rbinom(n, 1, p) - 1
y_1 <- which(y == 1)
y_0 <- which(y == -1)
x <- c()
x[y_1] <- rnorm(length(y_1), a, 0.1)
x[y_0] <- rnorm(length(y_0), b, 0.1)

sel <- cv.pmcid(x = x, y = y, delseq = deltaseq, k = 5,
               maxit = 100, tol = 1e-02)
dysel <- sel$'Selected delta'

result <- pmcid(x = x, y = y, n = n, delta = dysel,
               maxit = 100, tol = 1e-02, alpha = 0.05)
result$'Point estimate'
result$'Standard error'
result$'Confidence interval'
```

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