Pocket cards

At the following two pages you will find front and back of a pocket card for clinicians screening and randomising patients in the AID-ICU trial. There are four identical front pages and four identical back pages.

How to make the cards:

1. Write the personal username and password on the back side of the card
2. Print the two pages
3. Laminate the pages back to back
4. Cut out the cards



# Inclusion criteria

**Acute admission to the ICU**

**AND**

**Aged 18 years or above**

**AND**

**Diagnosed Delirium with validated screening tool: CAM-ICU (positive) or ICDSC ≥ 4.**

**Hotline: +45 9357 7750**

**Contact:** **aid-icu@cric.nu** **TURN OVER**

# Inclusion criteria

**Acute admission to the ICU**

**AND**

**Aged 18 years or above**

**AND**

**Diagnosed Delirium with validated screening tool: CAM-ICU (positive) or ICDSC ≥ 4.**

**Hotline: +45 9357 7750**

**Contact:** **aid-icu@cric.nu** **TURN OVER**

All patients should be screened for delirium two times daily. When a patient turns delirious please screen the patient for participation in the AID–ICU trial.

All patients should be screened for delirium two times daily. When a patient turns delirious please screen the patient for participation in the AID–ICU trial.

# Inclusion criteria

**Acute admission to the ICU**

**AND**

**Aged 18 years or above**

**AND**

**Diagnosed Delirium with validated screening tool: CAM-ICU (positive) or ICDSC ≥ 4.**

**Hotline: +45 9357 7750**

**Contact:** **aid-icu@cric.nu** **TURN OVER**

# Inclusion criteria

**Acute admission to the ICU**

**AND**

**Aged 18 years or above**

**AND**

**Diagnosed Delirium with validated screening tool: CAM-ICU (positive) or ICDSC ≥ 4.**

**Hotline: +45 9357 7750**

**Contact:** **aid-icu@cric.nu** **TURN OVER**



All patients should be screened for delirium two times daily. When a patient turns delirious please screen the patient for participation in the AID–ICU trial.

All patients should be screened for delirium two times daily. When a patient turns delirious please screen the patient for participation in the AID–ICU trial.

**1. Check inclusion criteria:**

See front page

**2. Go to** [**www.cric.nu/aid-icu**](http://www.cric.nu/aid-icu)

Follow the link (eCRF) and start screening

Check exclusion criteria when completing screening

**3. Prior to randomisation:**

* A negative pregnancy test (blood or urine) must be present in all fertile women (< 50 years)

**4. Follow procedure to randomise patient and to obtain the ampule identifier numbers**

**5. The identifier numbers of the ampules allocated to the patient will appear on the screen (each package contains three ampules)**

**6. Prescribe the trial medication to the patient, this include basic treatment 0,5ml three times daily and as needed trial medication 0,5ml maximum 5 additional doses daily.**

**7. Administer the medication to the patient**

**1. Check inclusion criteria:**

See front page

**2. Go to** [**www.cric.nu/aid-icu**](http://www.cric.nu/aid-icu)

Follow the link (eCRF) and start screening

Check exclusion criteria when completing screening

**3. Prior to randomisation:**

* A negative pregnancy test (blood or urine) must be present in all fertile women (< 50 years)

**4. Follow procedure to randomise patient and to obtain the ampule identifier numbers**

**5. The identifier numbers of the ampules allocated to the patient will appear on the screen (each package contains three ampules)**

**6. Prescribe the trial medication to the patient, this include basic treatment 0,5ml three times daily and as needed trial medication 0,5ml maximum 5 additional doses daily.**

**7. Administer the medication to the patient**:

**1. Check inclusion criteria:**

See front page

**2. Go to** [**www.cric.nu/aid-icu**](http://www.cric.nu/aid-icu)

Follow the link (eCRF) and start screening

Check exclusion criteria when completing screening

**3. Prior to randomisation:**

* A negative pregnancy test (blood or urine) must be present in all fertile women (< 50 years)

**4. Follow procedure to randomise patient and to obtain the ampule identifier numbers**

**5. The identifier numbers of the ampules allocated to the patient will appear on the screen (each package contains three ampules)**

**6. Prescribe the trial medication to the patient, this include basic treatment 0,5ml three times daily and as needed trial medication 0,5ml maximum 5 additional doses daily.**

**7. Administer the medication to the patient**

**1. Check inclusion criteria:**

See front page

**2. Go to** [**www.cric.nu/aid-icu**](http://www.cric.nu/aid-icu)

Follow the link (eCRF) and start screening

Check exclusion criteria when completing screening

**3. Prior to randomisation:**

* A negative pregnancy test (blood or urine) must be present in all fertile women (< 50 years)

**4. Follow procedure to randomise patient and to obtain the ampule identifier numbers**

**5. The identifier numbers of the ampules allocated to the patient will appear on the screen (each package contains three ampules)**

**6. Prescribe the trial medication to the patient, this include basic treatment 0,5ml three times daily and as needed trial medication 0,5ml maximum 5 additional doses daily.**

**7. Administer the medication to the patient**