

# Quality control for the FDG-PET studies in patients with disorders of consciousness

CSG PET workshop

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### Data quality checks - PET acquisition

- Confounding factors
  - Blood sugar / Glycemia
    - > Should be < 120
  - Arousal
    - > Patient should be awake
  - Least possible stimulation during glucose uptake
    - > Lights off, no stimulation unless sleeping
  - Neural inflammation might bias glucose uptake
- $\blacktriangleright$  Should be assured by following a proper protocol!  $\rightarrow$  Estelle

Boullaard et al., 2009

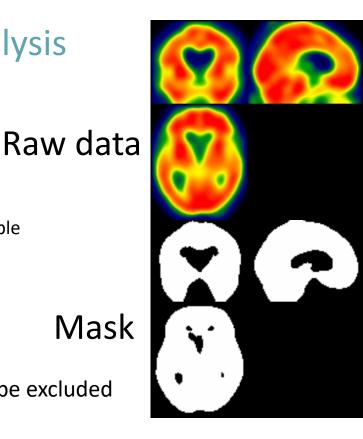


### Data quality checks – PET data analysis

- Single subject analysis
  - Healthy Control group
    - Larger is better (>20 subjects)
    - On the same scanner
  - Bad segmentation could bias SPM maps -
    - For single subject analysis it is sometimes unavoidable
      - Will underestimate hypometabolic areas

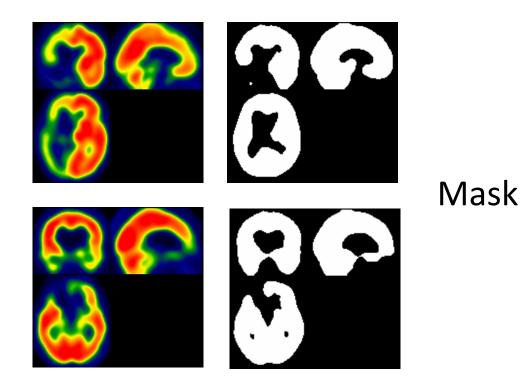
- Group analysis
  - Healthy Control group
  - Subjects with bad masks/segmentation should be excluded -

## Mask





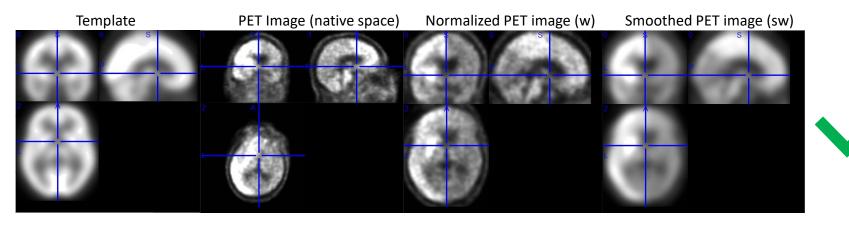
#### **Bad segmentation**



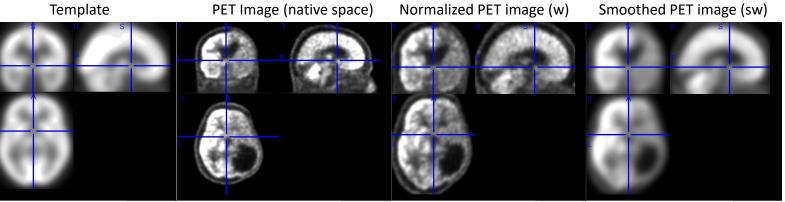
Raw data

#### **Bad normalization**





Template





#### **Bad normalization**





# Importance of good segmentation - group analysis

- ► After fitting your model evaluate:
  - 1. Mask image = voxels actually considered
    - > it should look like the brain volume
  - 2. RPV image = independent "resolution element"
    - > it should be relatively smooth in the brain tissue, except in some specific places, e.g. ventricles or brainstem
  - 3. the ESS image = unexplained variance in the model
    - > model should fit in the brain
- **Lesion/heterogeneous images negatively affect the GLM**, as the model may not be fitting there
  - Leads to: large extent with lots of unexplained signal and inflated smoothness estimates
- Consult experienced SPM user or developer for help.

Big thanks to Dr. Christophe Phillips & Dr. Mohamed Ali Bahri!



#### Thank you for your attention.

#### **Questions?**

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