# Diagnostic accuracy of neuroblastoma patient imaging with [18F]-mFBG PET-CT compared to [123I]mIBG scanning

No registrations found.

Ethical review	Positive opinion
Status	Recruitment stopped
Health condition type	-
Study type	Interventional

# **Summary**

### ID

NL-OMON28070

**Source** Nationaal Trial Register

**Brief title** MFBG PET-CT pilot study - neuroblastoma patient imaging -

#### Health condition

neuroblastoma

### **Sponsors and support**

**Primary sponsor:** Prinses Máxima Centrum for pediactric cancer **Source(s) of monetary or material Support:** KIKA

#### Intervention

### **Outcome measures**

#### **Primary outcome**

The primary study endpoints are the number of lesions and sites of disease detected with 18F-mFBG PET-CT compared to the current imaging standard of care, 123I-mIBG scan using the SIOPEN imaging

1 - Diagnostic accuracy of neuroblastoma patient imaging with [18F]-mFBG PET-CT comp ... 1-06-2025

scoring method for skeletal lesions and the total number of detected soft tissue lesions

#### Secondary outcome

o Determine optimum imaging time of 18F-mFBG PET CT(60 min vs 120-150 min post injection)

o Determine the estimation of radiation absorbed dose of 18F-mFBG

o Adverse events of 18F-mFBG injection and PET CT s

# **Study description**

#### **Background summary**

To compare [18F]mFBG PET-CT imaging for neuroblastoma patients with the current standard of imaging, [123I]mIBG SPECT, using the SIOPEN score for skeletal lesions and the number of detected soft tissue lesions as endpoints.

#### Study objective

[18F]mFBG PET-CT is probably better able to define neuroblastoma compared to the current imaging standard [123I] mIBG.

#### Study design

After 10 patients (6 months) and 20 patients (1 year)

#### Intervention

[18F]mFBG PET CT

# Contacts

**Public** Prinses Maxima Centrum Yvonne Ruchti

0889727272 **Scientific** Prinses Maxima Centrum Yvonne Ruchti

2 - Diagnostic accuracy of neuroblastoma patient imaging with [18F]-mFBG PET-CT comp ... 1-06-2025

# **Eligibility criteria**

# **Inclusion criteria**

patients with a (clinical suspicion of) neuroblastoma who are referred for [123I] mIBG imaging, 0-18 years of age

# **Exclusion criteria**

pregnancy of the patient and above 18 years old

# Study design

### Design

Study type:	Interventional
Intervention model:	Other
Allocation:	Non controlled trial
Masking:	Open (masking not used)
Control:	N/A , unknown

### Recruitment

NL	
Recruitment status:	Recruitment stopped
Start date (anticipated):	12-07-2020
Enrollment:	20
Туре:	Actual

### **IPD** sharing statement

Plan to share IPD: Undecided

# **Ethics review**

Positive opinionDate:12-11-2019Application type:First submission

# **Study registrations**

### Followed up by the following (possibly more current) registration

ID: 49035 Bron: ToetsingOnline Titel:

### Other (possibly less up-to-date) registrations in this register

No registrations found.

# In other registers

Register	ID
NTR-new	NL8152
ССМО	NL70903.041.19
OMON	NL-OMON49035

# **Study results**