# Comorbidities and treatment patterns among patients with acromegaly in Sweden – a register-linkage population-based study

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# Introduction

- Acromegaly is a rare disease most commonly caused by increased growth hormone (GH) secretion from a pituitary adenoma
- Acromegaly is associated with a number of comorbidities affecting e.g.
- the cardiovascular system, the musculoskeletal organs and metabolism
   Treatment of patients with acromegaly is complex, and includes surgery, pharmacotherapy and radiation therapy
- The objective was to describe comorbidities and treatment patterns among patients with acromegaly in Sweden

# Methods

# **Patients**

 Adult patients (aged > 18 years) with a first diagnosis of acromegaly due to a pituitary adenoma in Sweden between 1 July 2005 and 31 December 2013

## Design

- Observational, population-based study conducted in Sweden
- Linkage of nationwide registers with high precision via the unique personal identification number

### Data sources

- National Patient Register; for data on specialized health care contacts, including surgical and medical interventions
- Swedish Prescribed Drug Register; for data on prescribed medicines purchased in pharmacies
- Cause of Death Register; for data on time and cause of death (for censoring purposes)

# Endpoints and data analyses

- The presence of comorbidities was assessed
- The following treatment types were included: surgery, radiation therapy somatostatin analogues, dopamine agonists (DA) and growth hormone receptor antagonists (GHRA)
- The proportion of patients receiving each type of treatment at any time from date of diagnosis until death or the end of the study period was assessed, as well as the longitudinal treatment patterns
- The analyses were performed in the total study population as well as in
- subgroups based on time of diagnosis (before 2010 vs. on or after 2010)
- Descriptive statistics were used

# Results

# Patient population

- Predefined selection criteria identified 358 patients (48% men, mean age at diagnosis 50.0 [SD 15.3] years)
- 186 (52%) had received their first diagnosis before 2010, and 172 (48%) had been diagnosed on or after 2010
- The mean follow-up time from diagnosis to death or 31 December 2013 was 4.2 (SD 2.5) years
  - The follow-up time was 6.2 (SD 1.6) years among patients diagnosed before 2010, and 2.0 (SD 1.1) years among those diagnosed on or after 2010

### Comorbidities

- In the total study population, 81% (290/358) had at least one comorbidity

   Comorbidities were present among 88% (164/186) of patients
  - Comorbidities were present among 88% (164/186) of patients diagnosed before 2010 and among 73% (126/172) of patients diagnosed on or after 2010
- Figure 1 presents the 10 most common comorbidities

# Occurrence of each type of treatment

- The percentage of patients with each type of treatment is presented in Figure 2
- Two thirds (66%; 235/358) had undergone surgery at any time during follow-up
- 86 patients had more than one surgery
- One third (36%; 127/358) had been treated with SSA at any time during follow-up; most commonly as monotherapy (i.e. not in combination with DA or GHRA) (33%; 119/358)

## Longitudinal treatment patterns

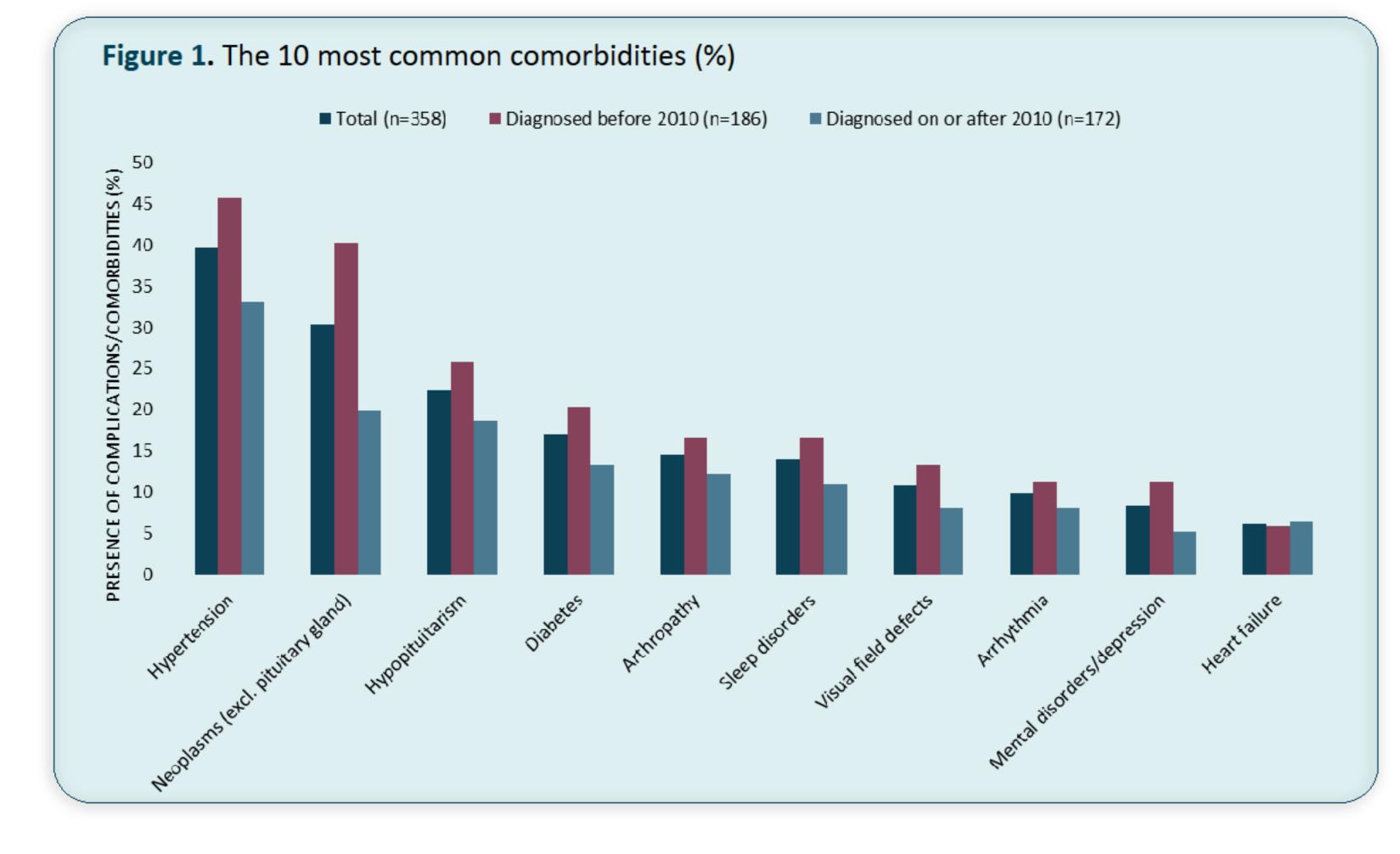
- First-line treatment was initiated on average 3.7 (mean; SD 6.9) months after diagnosis
- 16% (57/358) received no treatment for acromegaly during the follow-up
- Among the 301 patients who received treatment, the most common first-line treatments were surgery (n=180; 60%), SSA (n=64; 21%) and DA (n=41; 14%)
- Second-line treatments after primary surgical treatment are presented in Figure 3

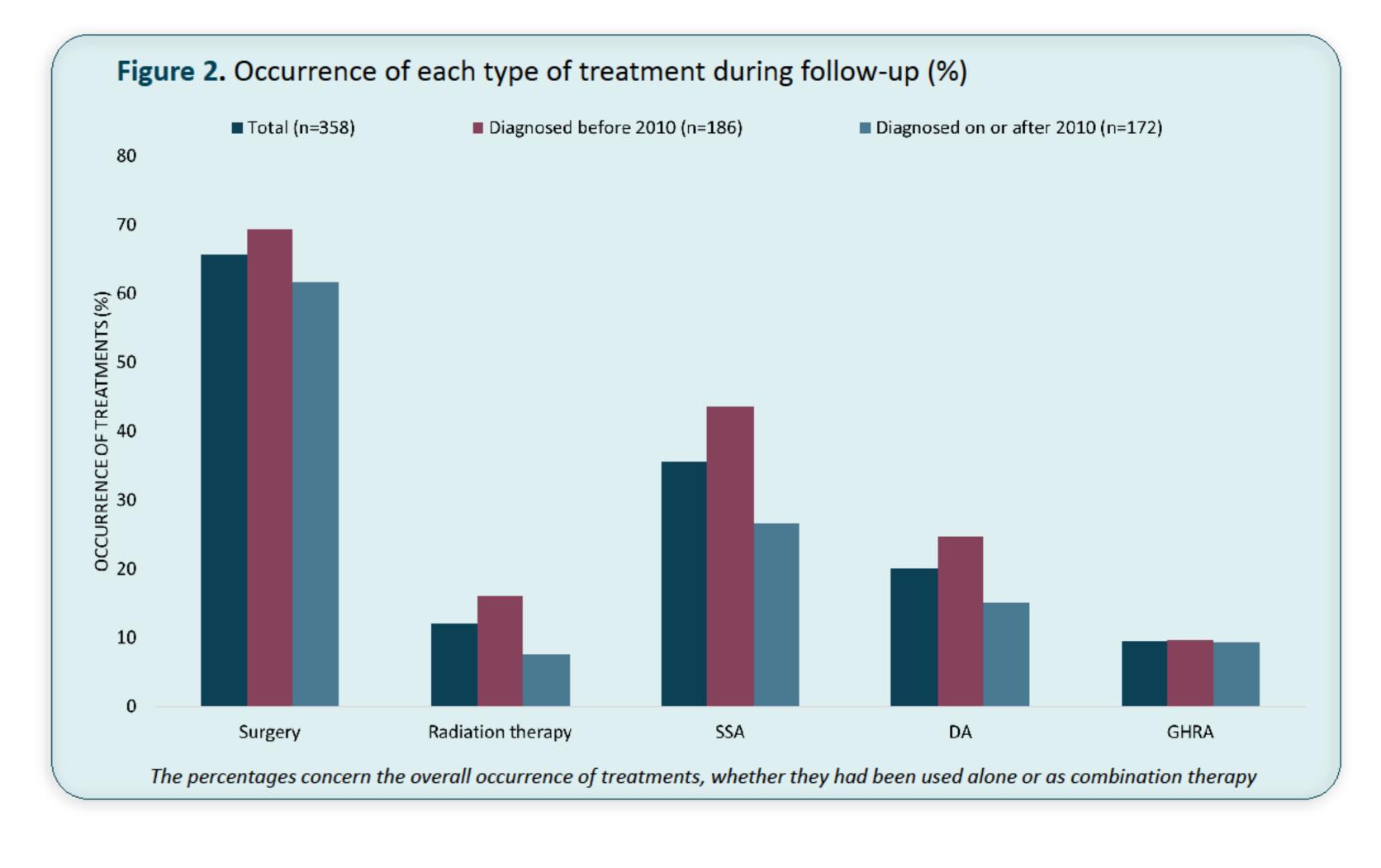
Time to treatment initiation was shorter among patients diagnosed on or

- Treatments following first-line SSA are presented in Figure 4
- after 2010 than among those diagnosed before 2010 (not shown in figure)

## Conclusions

- Comorbidity in patients with acromegaly is very high
- The most common first-line treatment was surgery, with the majority not receiving any second-line treatment
- SSA was the second most common first-line treatment, often followed by second-line surgery
- Time to treatment initiation was shorter among patients with a more recent diagnosis





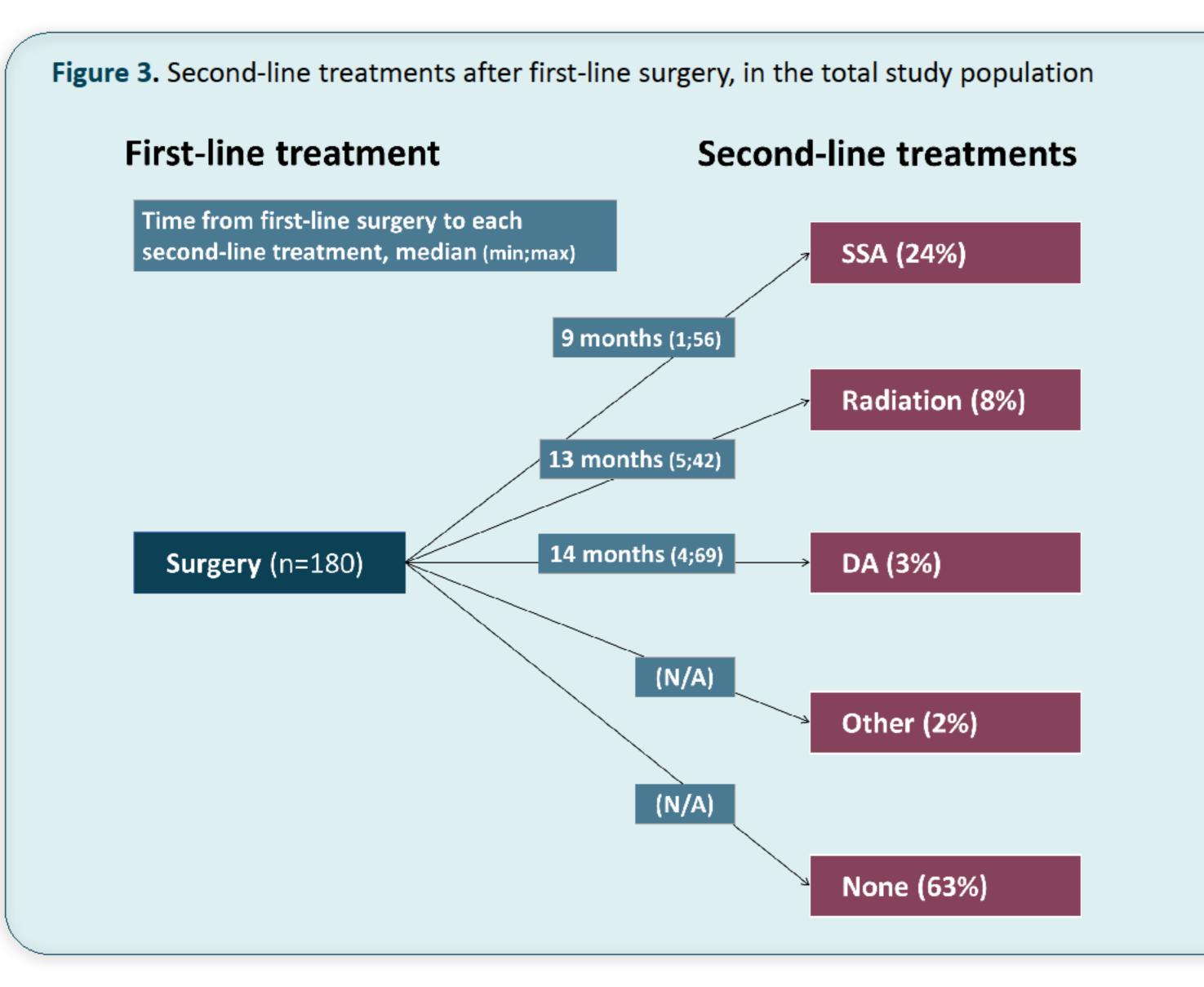


Figure 4. Second-line treatment

Second-line treatment

Time from first-line SSA to each second-line treatment, median (min;max)

Surgery (58%)

SSA + DA or GHRA (11%)

SSA (n=64)

(N/A)

Other (5%)

None (27%)

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