

# Incidence, Prevalence, and Risk Factors of Giant Cell Arteritis: A Targeted Literature Review

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## KEY FINDINGS & CONCLUSIONS

- GCA is more prevalent among individuals with Northern European descent compared to other ethnic groups
- Changes in healthcare policies and diagnostic coding practices may influence reported incidence and prevalence<sup>17</sup>
- Evidence gaps** identified from literature review:
  - Limited prevalence data** identified, with most results pertaining to incidence
  - Outdated data** as a lack of more recent publications identified
  - Heterogeneity** in study designs and diagnostic criteria: Limited ability to compare incidence and prevalence across geographies
- Further research from broad populations would be required to enhance understanding of the incidence and prevalence of GCA



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## INTRODUCTION

- Giant cell arteritis (GCA) is a systemic inflammatory disorder affecting medium and large arteries, typically in individuals aged >50 years<sup>1,2,3</sup>
- While glucocorticoids are the standard treatment, alternatives are necessary to reduce side effects and sustain remission<sup>4,5</sup>
- Substantial overlap exists between GCA and polymyalgia rheumatica (PMR)<sup>6</sup>
- Understanding the evolving epidemiology of GCA is essential for assessing disease burden and developing effective treatments

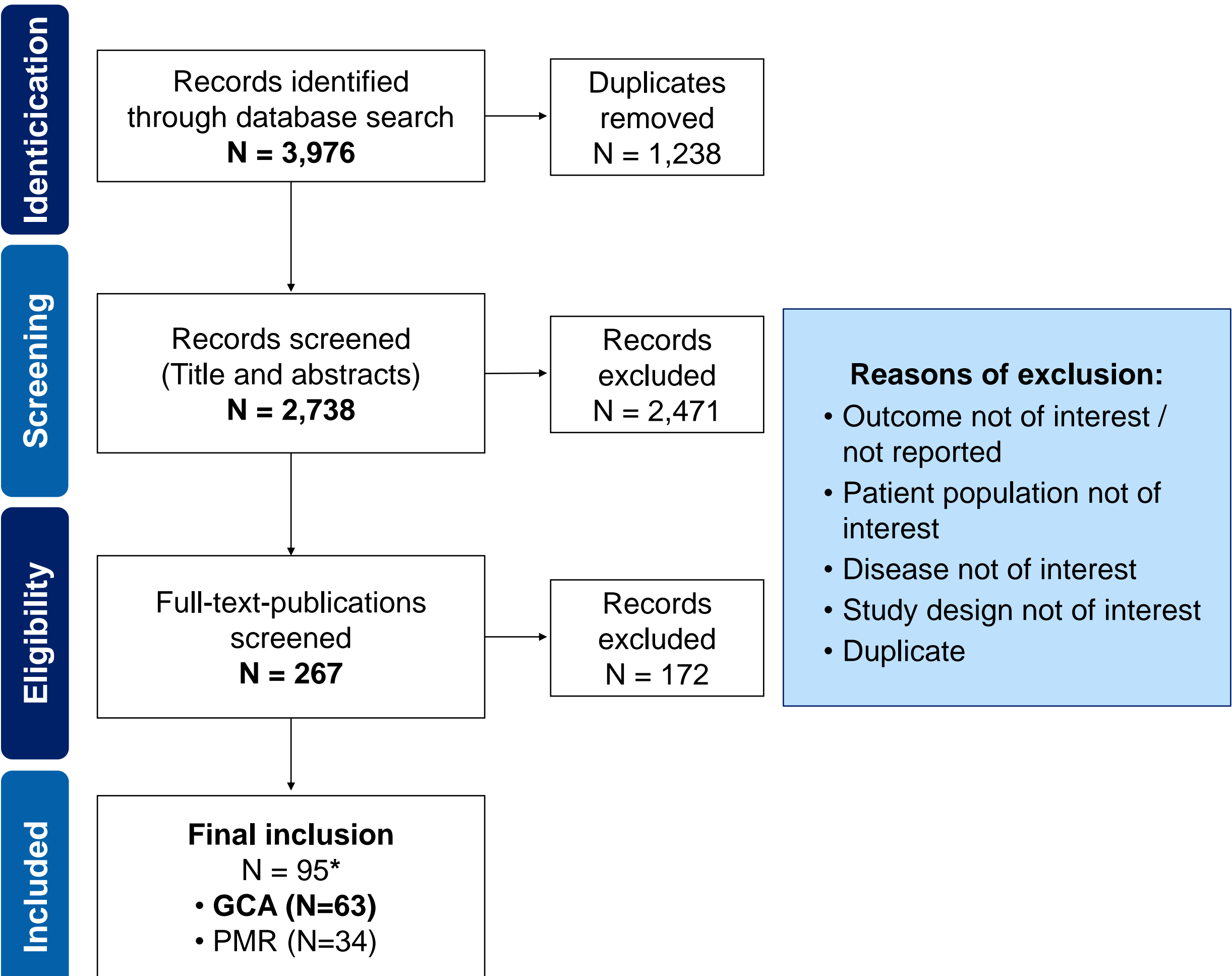
## OBJECTIVES

- To identify and summarize evidence from studies on the epidemiology of GCA focusing on prevalence, incidence, and risk factors

## RESULTS

- A total of 3,976 records were identified, of which 95 were included and 63 specifically related to GCA (**Figure 2**)

Figure 2. PRISMA Flow Chart<sup>#</sup>



\*Note: The final included publications (n=95) encompass publications for both GCA and PMR indications  
Refer to poster EPH111 details on studies relating to PMR  
GCA, Giant Cell Arteritis; PMR, Polymyalgia Rheumatica; PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analyses; SLRs, Systematic Literature Reviews. <sup>#</sup>Page et al. BMJ 2021;372:n71 – Applied to a targeted literature review process

Table 1. Prevalence of GCA reported across key geographies

Country (Region)	Type	Prevalence per 100,000 (95% CI) (aged ≥50 years)		
		Overall	Male	Female
USA <sup>7</sup> (Olmsted County, Minnesota)	Point	<b>204</b> (161-254)	<b>91</b> (46-156)	<b>304</b> (229-375)
*Germany <sup>8</sup> (North)	Period	<b>44.0</b> (39.9-48.1)	<b>21.9</b> (19.0-24.8)	<b>61.2</b> (56.4-66.0)
*UK <sup>9</sup> (Norfolk)	Cumulative	<b>410</b> (230-580)	<b>290</b> (70-500)	<b>520</b> (250-790)
Spain <sup>10</sup> (Costa del Sol)	Period	<b>12.2</b> (5.6-18.9)	NR	NR
Italy <sup>11</sup> (Reggio Emilia)	Point	<b>101.3</b> (88.4-115.4)	<b>22.6</b> (17.2-29.1)	<b>60.5</b> (51.6-70.5)

CI: Confidence Interval; GCA: Giant Cell Arteritis; NR: Not Reported; UK: United Kingdom; USA: United States of America  
\*Converted to equivalent per 100,000 population; \*GP diagnoses; aged ≥55 years

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## METHODS

- A comprehensive targeted literature review was performed with predefined PICOS criteria in February 2024 (**Figure 1**)
- Strategies were applied across various databases, including Embase®, Medline®, CDSR, CENTRAL, DARE, and HTA, along with manual searches on congress websites and bibliographic sources
- All records identified were screened during the first (title/abstract) and second (full text) stages
- Screening and data extractions were done by one reviewer, with uncertainties discussed and quality checks performed by a second independent reviewer
- Study selection, data extraction and reporting adhered to current best practices
- Only studies published in English language were considered for inclusion

### Prevalence

- Global pooled prevalence of GCA was estimated as **51.74/100,00 people** (95% CI: 42.04, 61.43) aged >50 years<sup>3</sup>
- Variability was observed across country-specific studies, with women showing ~2-3 times higher prevalence rates (**Table 1**)<sup>7-9,11</sup>

### Incidence

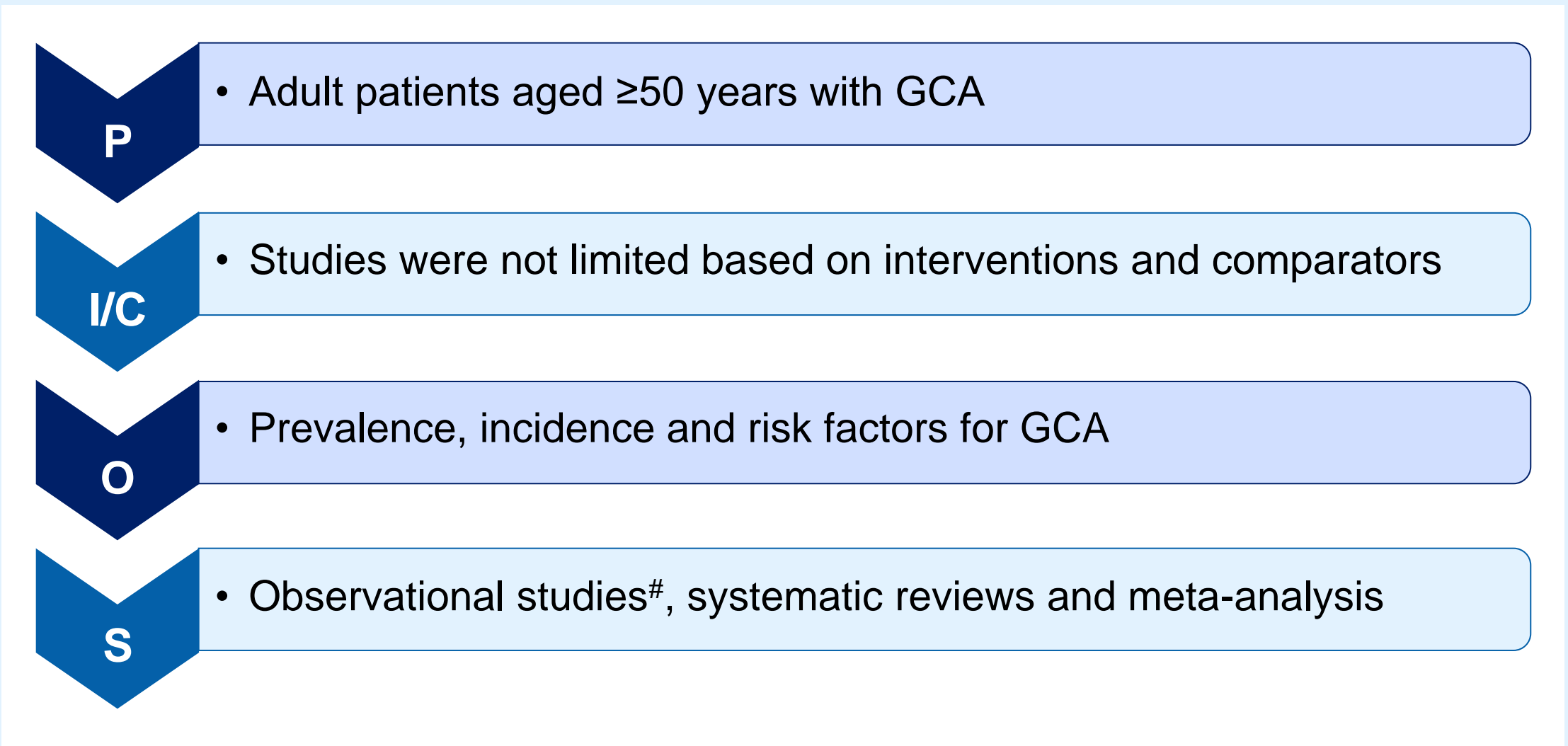
- A recent systematic literature review and Network Meta-analysis estimated the pooled global (**10 (9.22-10.78) per 100,000**) and regional incidence of GCA among people aged >50 years<sup>3</sup>
- Pooled GCA incidence showed **nearly 3-fold variation between regions** and was reported to **be highest in Scandinavia**<sup>3</sup>
- The included regions from highest to lowest incidence per 100,000 persons (95% CI) aged >50 years were:<sup>3</sup>
  - Scandinavia: **21.57** (18.90-24.23)
  - North and South America: **10.89** (8.78-13.00)
  - Oceania: **7.85** (-1.48-17.19)
  - Europe: **7.26** (6.05-8.47)
  - Middle East: **5.73** (4.20-7.26)
  - Africa: **4.62** (0.05-9.20)
  - East Asia: **0.34** (0.12-0.56)
- Country-specific, with higher incidence in females (**Table 2**)

Table 2. Incidence of GCA reported across key countries

Country (Region)	Incidence per 100,000 (95% CI) (aged ≥50 years)		
	Overall	Male	Female
USA <sup>12</sup> (Olmsted County, Minnesota)	<b>19.8</b> (15.2-24.3)	<b>10.1</b> (5.0-15.3)	<b>27.0</b> (20.0-33.9)
Canada <sup>13</sup> (Ontario)	<b>4.9</b> (4.2–5.6)	NR	NR
UK <sup>14</sup> (Norfolk)	<b>9.8</b> (8.6-11.2)	<b>7.0</b> (5.6-8.7)	<b>12.4</b> (10.6-14.5)
France <sup>15</sup> (Paris)	<b>7.6</b> (5.9–9.8)	<b>5.4</b> (3.3–8.3)	<b>9.5</b> (6.9–12.9)
Italy <sup>11</sup> (Reggio Emilia)	<b>8.3</b> (7.1-9.4)	<b>5.3</b> (3.9 - 6.6)	<b>10.8</b> (9.0 - 12.6)
China <sup>16</sup> (Hong Kong)	<b>0.34</b> (NR)	NR	NR

CI: Confidence Interval; GCA: Giant Cell Arteritis; NR: Not Reported; UK: United Kingdom; USA: United States of America

Figure 1. PICOS Criteria\*



\*Prospective, retrospective, cohort, cross-sectional, case-control, claims database, registry studies.  
C: Comparators; GCA: Giant Cell Arteritis; I: Intervention; O: Outcomes;; P: Patient population; S: Study design  
\*Methley et al. BMC Health Services Research (2014) 14:579

### Risk Factors

- Several risk factors associated with GCA were identified and can be broadly categorized into non-modifiable and modifiable, as summarized in **Figures 3** and **4**

Figure 3. Non-modifiable risk factors

<b>Age</b>	<b>Geography</b>
Increased risk with <b>age</b> : <ul style="list-style-type: none"><li><b>Rare</b> in people <b>&lt;50 years</b><sup>3,17</sup></li><li><b>Most common</b> in people <b>≥75 years</b><sup>12</sup></li></ul>	<b>Incidence in Scandinavia</b> <sup>*3,17</sup> : <ul style="list-style-type: none"><li><b>3 times higher</b> than in the <b>rest of Europe</b></li><li><b>6 times higher</b> than in <b>East Asia</b></li></ul>
<b>Female Gender</b>	<b>PMR Presence</b>
GCA is <b>2.5-3 times</b> more likely in <b>women</b> than <b>men</b> <sup>3,17</sup>	Approximately <b>22%</b> of those at the time of <b>PMR diagnosis</b> have <b>GCA symptoms</b> <sup>6</sup>

\*Communities in the USA with Scandinavian ancestry are also subject to higher rates  
GCA: Giant Cell Arteritis; PMR: Polymyalgia Rheumatica

Figure 4. Modifiable risk factors

<b>Smoking</b>	<b>Biomarkers</b>
Increased risk vs. non-smokers <sup>18</sup> : <ul style="list-style-type: none"><li><b>Current smokers</b> (OR: 1.18, 95% CI: 1.01-1.38)</li><li><b>Former smokers</b> (OR: 1.19, 95% CI: 1.01-1.39)</li></ul>	Increased risk associated with <b>higher levels of biomarkers</b> <sup>19</sup> : <ul style="list-style-type: none"><li><b>IFN-γ</b> (OR: 2.37, 95% CI: 1.14-4.92)</li><li><b>MCP3</b> (OR: 3.74, 95% CI: 1.26-11.07)</li></ul>
<b>Cardiovascular Factors</b>	<b>BMI</b>
Significant increased risk associated with <sup>20</sup> : <ul style="list-style-type: none"><li><b>Hyperlipidemia</b> (SHR: 1.27; 95% CI: 1.12–1.42; p&lt;0.01)</li><li><b>Hypertension</b> (SHR: 1.22; 95% CI: 1.09-1.36; p&lt;0.05)</li></ul>	Statistically significant <b>inverse correlation</b> : → <b>8% risk increase</b> per 1.0 kg/m <sup>2</sup> <b>BMI decrease</b> <sup>21</sup>
<b>Infections</b>	<b>Gout</b>
Increased risk with <sup>22,23</sup> : <ul style="list-style-type: none"><li><b>Acute upper respiratory tract infection</b> (OR: 1.77, 95% CI: 1.47–2.14)</li><li><b>Influenza</b> and <b>pneumonia</b> infections (OR: 1.72, 95%CI: 1.35–2.19)</li><li><b>Prior herpes zoster</b> infections (OR: 1.20, 95% CI: 1.08-1.32)</li><li><b>Prior overall infections</b> (OR: 1.27, 95% CI: 1.18-1.37)</li></ul>	<b>Gout</b> associated with <b>&gt;2 times higher risk of developing GCA in older adults</b> , independent of known GCA risk factors (HR: 2.05, 95% CI: 1.76-2.40) <sup>24</sup>
	<b>ARBs</b>
	<b>Incidence higher in initiators of ARBs</b> compared to <b>ACEis</b> . Increased hazard ratio for ARB initiators (HR: 1.55; 95% CI: 1.16-2.06) <sup>25</sup>

ACEi: Angiotensin-Converting Enzyme Inhibitors; ARB: Angiotensin receptor blockers; BMI: Body mass index; CI: Confidence Interval; HR: Hazards Ratio; GCA: Giant Cell Arteritis; IFN-γ: Interferon Gamma; MCP3: Monocyte chemotactic protein 3; OR: Odds Ratio; PMR: Polymyalgia Rheumatica; SHR: Sub-Hazard Ratio

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## Disclosures

The authors wish to disclose the following: Sarah Jane McKenna and Jessica Commane are current employees of Novartis Ireland Ltd. and shareholders with Novartis and Sandoz. Samprati Avasthi and Ramakrishna GS are current employees of Novartis Healthcare Pvt. Ltd., Hyderabad, India