

Identification and Description of Real-World Data Sources for Giant Cell Arteritis (GCA) and Polymyalgia Rheumatica (PMR)

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KEY FINDINGS & CONCLUSIONS

- There is a lack of disease-specific sources for both GCA and PMR, with only 9 and 1 registries identified, respectively.
- This underscores the need for future collaborations to collect robust RWD to better understand GCA and PMR.
- Limitations: Description of the identified RWD was based on information publicly available. Data owners were not contacted to confirm the accuracy of this evidence.

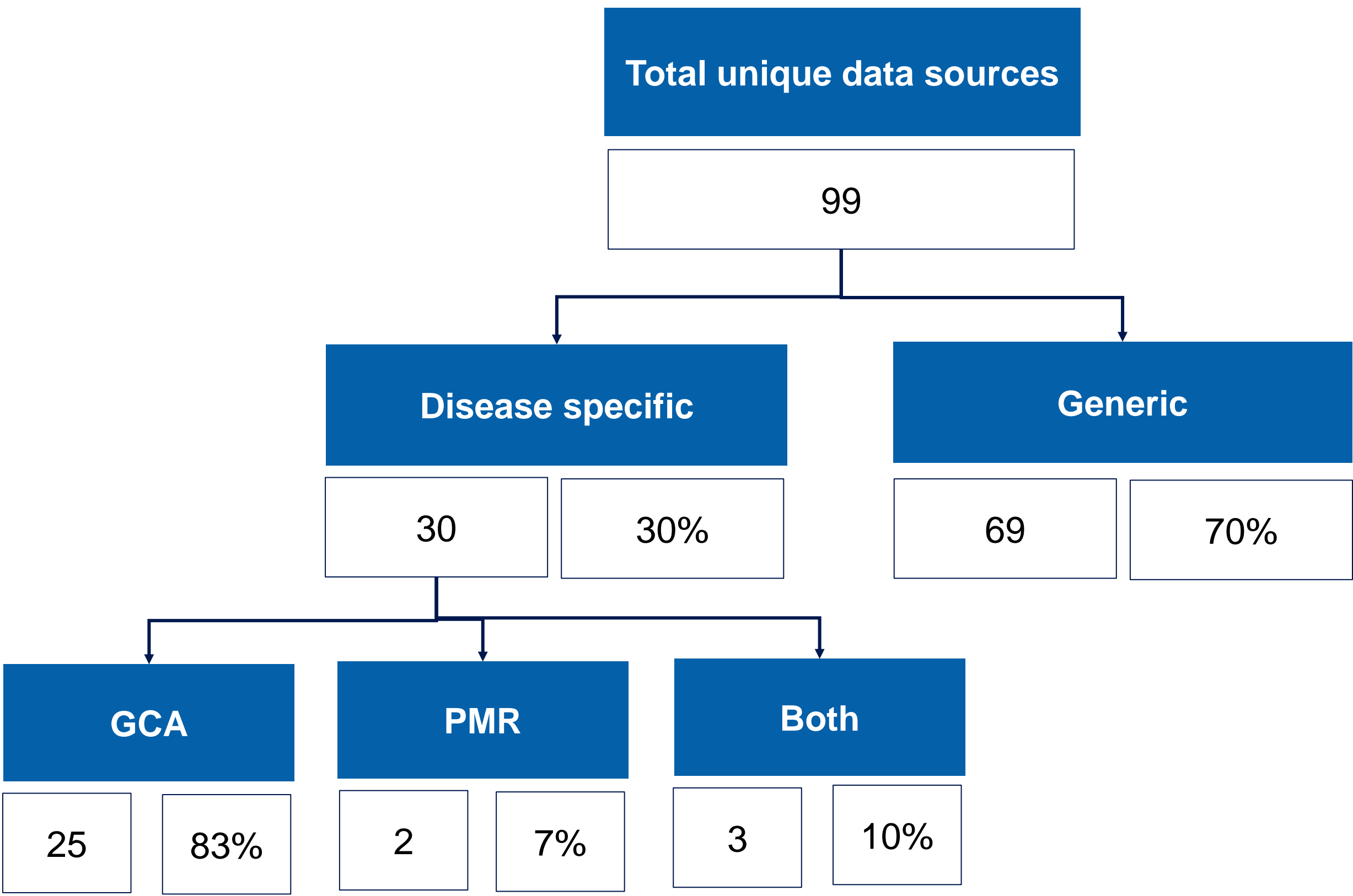
INTRODUCTION

- GCA is a chronic, idiopathic vasculitis of large and medium arteries, typically in individuals aged >50 years, resulting in chronic inflammation, which leads to vascular remodeling and occlusion.^{1,2,3} GCA can lead to serious complications such as permanent visual loss and stroke, if left untreated.
- PMR is an inflammatory rheumatic disorder characterized by muscle pain and stiffness, especially in the neck, shoulder, and pelvic girdle⁴
- 16% - 21% of PMR patients⁵ can develop GCA while 50% of GCA patients⁶ report PMR as a comorbidity

RESULTS

- A total of 2242 publications were retrieved from the literature search and 99 unique data sources were identified.
- Detailed information for up to 44 variables was extracted for each unique data source, when provided in the publications.
- Amongst all the unique data sources, only one-third of them were disease-specific and most (>80%) were focused on GCA.

Figure 1. GCA and PMR data sources classification



- Across all sources (n=99), the most widely reported parameters were age (95%), gender (93%), labs including C-reactive protein (62%) and erythrocyte sedimentation rate (48%), temporal artery biopsy (48%), glucocorticoids (60%), visual complications (46%), certain comorbidities like diabetes mellitus (44%).
- The least reported parameters include genetics (2%), upper limb claudication (3%), leflunomide (4%), emergency visits and costs (3%), and PROs (1-3%).
- Detailed completion rate of variables across the different categories is given in Figure 2.

GCA: Giant Cell Arteritis, PMR: Polymyalgia Rheumatica, PRO: Patient Reported Outcomes

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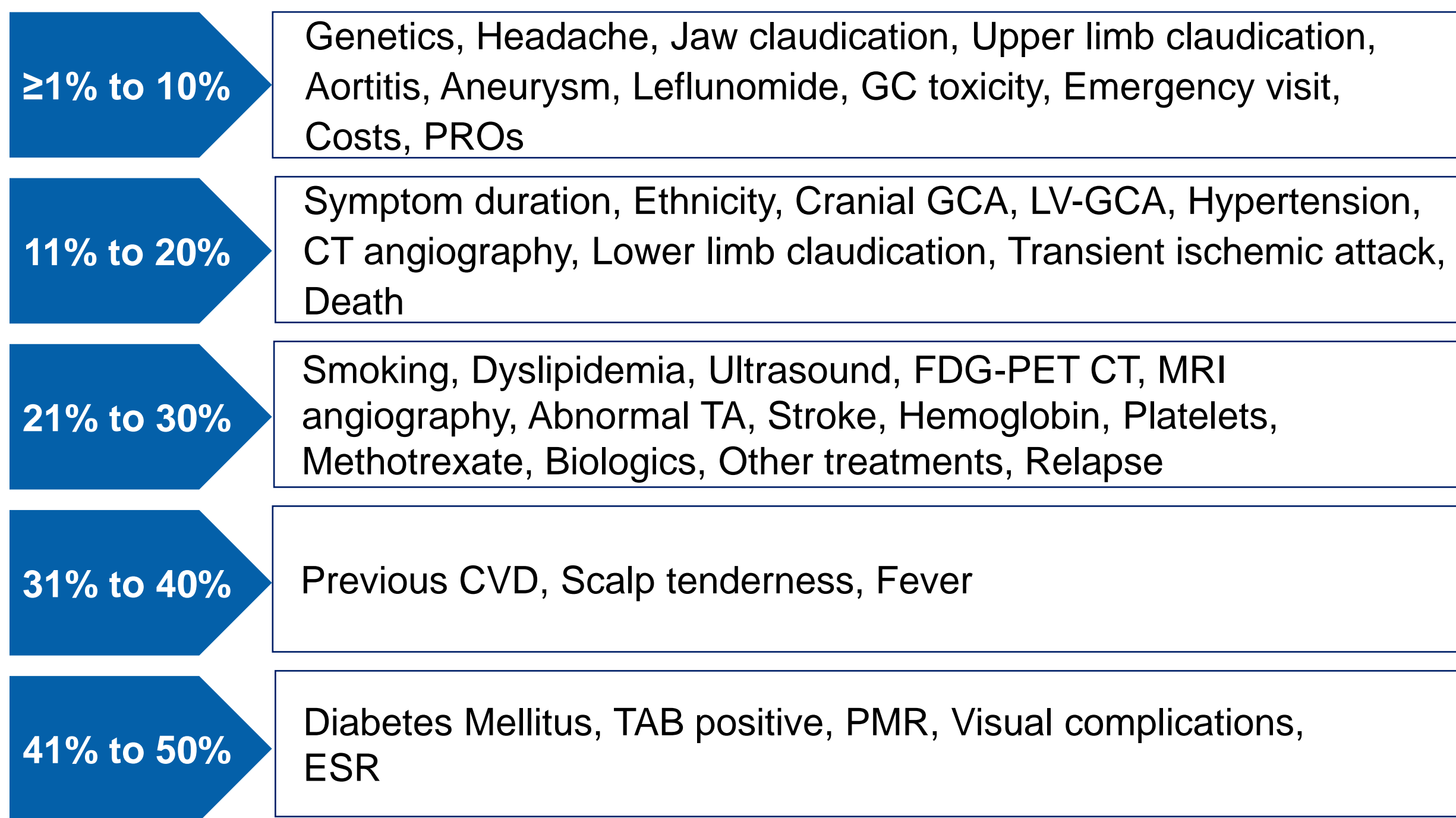
OBJECTIVES

- Given the growing interest for real-world data (RWD), we aimed to identify disease-specific RWD for Giant Cell Arteritis (GCA) and Polymyalgia Rheumatica (PMR) and to describe their key characteristics (variables collected, geographical scope, design etc.)

METHODS

Database search	EMBASE / Medline (2019 - May 2024; past 5 years) based on defined search terms and PICO framework
Screening	• Title / abstract screening • Full text screening (for articles included in 1)
Linking and Finalization	• Link publications based on unique data sources • Finalize unique list of data sources
Data mapping	• Identify and map different variables using data abstraction tool

Figure 2. Completion rate of selected variables



GC: Glucocorticoid, PRO: Patient Reported Outcomes, LV-GCA: Large Vessel Giant Cell Arteritis, CT: Computed Tomography, FDG-PET: Fluorodeoxyglucose Positron Emission Tomography, MRI: Magnetic Resonance Imaging, TA: Temporal artery, CVD: Cardiovascular Disease, TAB: Temporal Artery Biopsy, ESR: Erythrocyte Sedimentation Rate

- The top three variables which were the most reported across each of the categories are indicated in Figure 3.

Figure 3. Top 3 variables across the different categories

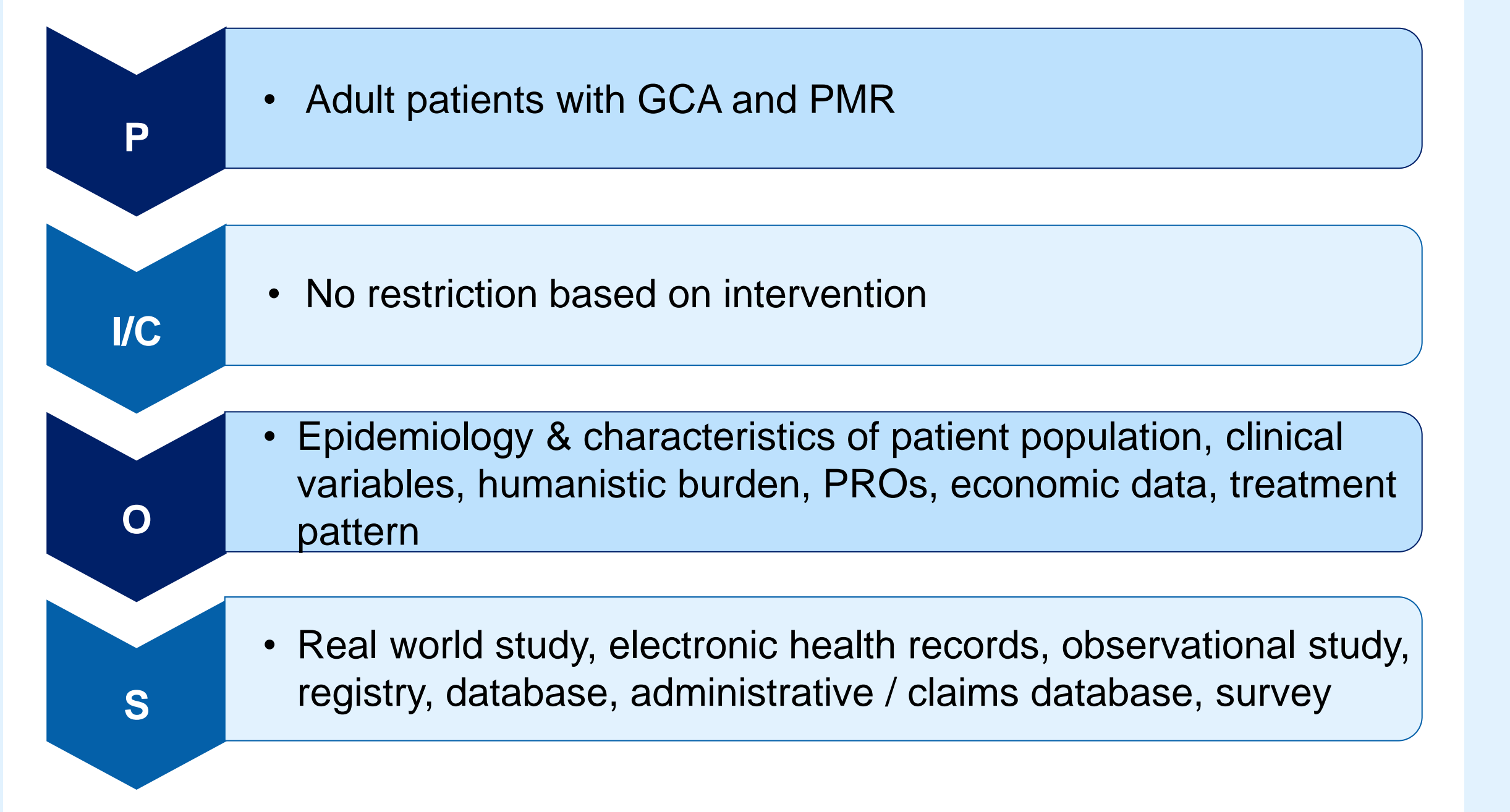
Demographics and diagnosis	• Age (95%) • Gender (93%) • Temporal artery biopsy (48%)
Risk factors	• Diabetes Mellitus (44%) • Hypertension (40%) • Previous CVD (31%)
Clinical outcomes	• Occurrence of PMR (48%) • Visual Complications (46%) • Scalp Tenderness (34%)
HRCU	• Hospitalizations (10%) • Costs (3%) • Visits (3%)
Treatments	• Glucocorticoids (60%) • Biologics (25%) • Other drug classes (25%)

CVD: Cardiovascular disease, PMR: Polymyalgia Rheumatica, HRCU: Healthcare Resource Utilization

References

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Figure 1. PICOS Criteria



P: Population, I: Intervention, C: Comparator, O: Outcomes, S: Study Design

- Amongst the 30 disease specific data sources, there were 10 registries, of this, 9 were GCA related and 1 was PMR.
- Majority of these disease specific data sources were multi-center (69%).
- In terms of geographic coverage, Europe (83%) was largely covered, followed by North America (13%) and Japan (3%).

Table 1. Disease specific registries

Data Source	Population	Geography	N	Years covered	Center
Diagnostic and Classification Criteria in Vasculitis (DCVAS)	GCA	International	6991	2011-17	Multicenter
International Collaboration - GCA	GCA	Europe	778	2015	Multicenter
HAS-GCA Study	GCA	Europe	229	2019-22	Multicenter
Vasculitis Clinical Research Consortium (VCRC) Study	GCA	North America	426	2006-19	Multicenter
ARTEMIS study	GCA	France	306	2018 (Aug-Nov)	Multicenter
The Joint Vasculitis Registry in German-speaking countries (GeVas)	GCA	Germany, Austria, Switzerland	517	2019-ongoing	Multicenter
Real-Life Treatment and Safety - GCA Cohort Study (REATS)	GCA	Germany	395	Unknown	Multicenter
Spanish Giant Cell Arteritis Registry (ARTESER)	GCA	Spain	1636	2013-19	Multicenter
Rheumatology Informatics System for Effectiveness (RISE)	PMR	USA	240,000	2016-22	Multicenter

HAS-GCA: Halo and SGCAPS Score for Giant Cell Arteritis

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Disclosures

The authors wish to disclose the following: Julie Le Moal Mouchet and Valeria Jordan Mondragon are current employees of Novartis Pharma AG and Novartis Pharmaceuticals Corporation and shareholders with Novartis. Samprati Avasthi and Jagrut Vaishnav are current employees of Novartis Healthcare Pvt. Ltd., Hyderabad, India