## Classification of Patients with Cardiovascular Diseases: Data from the Ontario Best Practices Research Initiative (OBRI) Kangping Cui<sup>1,2</sup>, Mohammad Movahedi<sup>1,</sup>, Claire Bombardier<sup>1,3,4</sup>, Bindee Kuriya<sup>1,4</sup> and OBRI Investigators

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## BACKGROUND

- Cardiovascular disease (CVD) is a major comorbidity and a leading cause of death among rheumatoid arthritis (RA) patients<sup>1-2</sup>.
- There are limited data on the prevalence and characteristics of RA patients with CVD in Canada.
- Ongoing research exploring CVD and its effect on RA disease outcomes has been undertaken at the Ontario Best-practices Research Initiative Rheumatoid Arthritis (OBRI-RA) Registry.
- Within the Registry, physician-reported cardiovascular disease has a broad definition, some of which do not meet the strict definition of "cardiovascular" disease".
- Precisely identify and classify patients with cardiovascular disease and its risk factors is imperative for the success of future studies.

## **OBJECTIVE**

To develop an algorithm in identifying and confirming the diagnosis of patients meeting the definition of CVD and CVD risk factors.

## METHODS

- Data (as of Jan 2017) were collected from the OBRI, a clinical registry adult RA patients followed in routine care in Ontario, Canada.
- CVD is defined as having one or more of the following:
  - Prior myocardial infarction (MI)
  - Interventions for coronary artery disease (CAD)
  - Transient ischemic attack (TIA)
  - Stroke
  - Peripheral arterial disease (PAD)
- CVD risk factors used in this study includes the presence of:
  - Hypertension (HTN)
  - Dyslipidemia (DLD)
  - Diabetes Mellitus (DM)
  - Being a current smoker at the OBRI enrolment
- The classification of CVD risk factors and CVD are outlined in **Figure 1 and 2** respectively.



### Fig. 1. Classification of DLD (as example for CVD risk factors classification) with physician and patient reported information.

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### Fig. 2. Classification of CVD with physician and patient reported information.

\* MI: myocardial infarction; CAD: coronary artery disease; PAD: peripheral artery disease, CVA: cerebral vascular accident (including stroke and TIA). \*\* Clinical judgement is applied at this step. Information considered includes whether the patient is on anti-platelet agents, anti-hypertensives, cholesterol control medications, or nitroglycerines. For uncertain cases, a second reviewer's opinion is weighed in.

### RESULTS



Fig. 3. The prevalence of CVD and CVD RF.

### Table 1. The prevalence of individual CVD and CVD RF.

Individual CVD	N (%)
MI/CAD	98 (4.8%)
Stroke/TIA	11 (0.5%)
PAD	1 (0.1%)
Individual CVD risk factor	N (%)
Hypertension	670 (33.0%)
Diabetes Mellitus	165 (8.1%)
Dyslipidemia	401 (19.7%)
Current smoking	346 (17.0%)

## RESULTS

### Table 2. Comparison of prevalence of CVD and CVD RF in OBRI-RA cohort against selected international cohorts.

	OBRI	US pharMetrics*	COMORA**	Swedish RA Registry <sup>#</sup>
CVD	5.4%	4.0%-8.8%	6%	4.5% (MI only)
HTN	33.0%	31%	40%	27.3%
DLD	19.7%	28.3%	32%	n/a
DM	8.1%	10.4%	14%	8.1%
Smoking	17.0%	n/a	20%	29.8%

\* Largest integrated US health plan database with 28,000 RA patients \*\* International registry with 4500+ patients recruited in 17 participating countries from Asia, Europe, and America. <sup>#</sup> Early RA registry based in Sweden.

# CONCLUSIONS

- cardiovascular disease.
- the OBRI-RA registry.

- outcomes.

# REFERENCES

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Additional patients were classified after medication review: 17 subjects (15.7%) for CVD, 207 for HTN (31%), 291 for DLD (73%), and 22 for DM (13%). Chart review of 55 patients showed sensitivity of 100% for CVD, 78% for HTN, and 42% for DLD classification.

• 110 out of 2033 patients (5.4%) was classified as having CVD (Fig. 3).

The prevalence of CVD, HTN and DM in OBRI-RA cohort was comparable to international cohorts, while the prevalence of DLD and smoking was lower (Table 2). Among patients with CVD RFs, the majority has one risk factor (36.7%), followed by two risk factors (11.7%).

We have successfully developed an algorithm for classifying patients with

The algorithm is applicable for the classification of other comorbidities captured in

This study highlights the complexity of data extraction from clinical registry.

The lower prevalence of DLD could be explained by the local-regional difference in disease prevalence as well as potential classification bias.

Further analysis is underway to explore the effects of CVD and its RF on RA

1. Choy E et al. Cardiovascular risk in rheumatoid arthritis: recent advances in the understanding of the pivotal role of inflammation, risk predictors and the impact of treatment. Rheumatology (Oxford, England) Published Online First: 2014.

2. Peters MJ et al. EULAR evidence-based recommendations for cardiovascular risk management in patients with rheumatoid arthritis and other forms of inflammatory arthritis. Annals of the rheumatic diseases 2010;69:325–31.

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