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Psoriasis is an independent predictor of increased risk of allergic reaction during percutaneous coronary interventions. Big data analysis from the Polish National PCI Registry (ORPKI)

Running title: Psoriasis and complications during PCI

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Abstract

Background: The presence of psoriasis is currently considered by the European Society of Cardiology (ESC) cardiovascular prevention guidelines of 2016 as one possible cardiovascular risk factor. Patients with psoriasis and concomitant coronary artery disease treated by means of percutaneous coronary intervention (PCI) are a fairly large subgroup of patients that have been usually omitted in mainstream research. The aim herein, was to identify the incidence of psoriasis, baseline characteristics and periprocedural outcome with a special focus on procedural complications in patients undergoing percutaneous coronary procedures.

Methods: All consecutive patients who had either coronary angiography or coronary angiography with immediate PCI in Poland in 2014 and 2015 were included. Patients were
assigned to two groups based on previous diagnosis: with psoriasis and without psoriasis. Clinical outcome was defined as any periprocedural death.

**Results:** There were 405,078 patients included in this analysis. Psoriasis (moderate or severe) was diagnosed in 1507 (0.4%) of them. Psoriasis was an independent predictor of allergic reaction occurrence (odds ratio [OR] 6.02; 95% confidence interval [CI] 1.44–25.22; p = 0.014). After propensity score adjustment, psoriasis remained a significant predictor of allergic reaction (OR 5, 95% CI 1.2–20.7; p = 0.0245). There were no differences in rates of periprocedural deaths in patients with or without psoriasis (death: 0.95% vs. 0.62%, p > 0.05).

**Conclusions:** Severe or moderate psoriasis is an independent risk factor for the occurrence of allergic reaction during percutaneous coronary procedures. There were no differences in periprocedural mortality and complications in patients with versus those without psoriasis.

**Key words:** infarction, coronary, psoriasis, anaphylaxis

**Introduction**

Psoriasis is a chronic autoimmune inflammatory disease, the prevalence of which ranges from 1.5% to 4.7% in the general population making it a significant healthcare issue [1–3]. The immune-mediated inflammatory pathogenesis of psoriasis is linked to another inflammatory disease which is atherosclerosis [4, 5]. The presence of psoriasis is currently considered by the European Society of Cardiology (ESC) cardiovascular prevention guidelines of 2016 as one possible cardiovascular risk factors within a wide group of autoimmune and inflammatory diseases [6]. Currently the American Heart Association (AHA) recommendations include risk factor screening in patients with psoriasis from as early as age 20 and by the age 40 a full list of these including various cardio-metabolic parameters need to be identified [7, 29]. Retrospective data concerning the effect of psoriasis on cardiovascular morbidity and mortality seem to confirm its adverse outcome on the occurrence of myocardial infarction or development of coronary artery disease, however, meta-analysis and literature reviews provide inconclusive data, probably because of confounding factors and the moderate quality of cardiovascular risk factor reporting (psoriasis) in large datasets [8–12]. In the Rotterdam study, authors did not find a correlation
between mild psoriasis and risk for the development of cardiovascular events or atherosclerosis which seems to support the general consensus that if any, psoriasis may have its deleterious effect only if at least moderate or severe presentation is expressed [13]. A combined clinical endpoint of death, myocardial infarction and stroke was more frequent in patients with severe psoriasis than in those without, which is the only prospective observation available thus far [14, 15].

Patients with psoriasis and concomitant coronary artery disease treated by means of percutaneous coronary intervention (PCI) are a fairly large (in definite numbers) subgroup of patients that have usually been omitted in mainstream research.

The main aim of this analysis was to identify the incidence of psoriasis, baseline characteristics and periprocedural outcome with a special focus on procedural complications in this subgroup of patients.

Methods

The Polish National PCI database (ORPKI) is a registry (with eCRF) for all percutaneous coronary procedures that have been performed in Poland since 2004 and is currently operated by the Jagiellonian University Medical College in Krakow. Annual reports are published each year and the registry rationale has been described before [16, 17]. In this manuscript, data for the calendar year 2014 (January–December) and 2015 (January–December) were analyzed. All consecutive patients who had either coronary angiography or coronary angiography with immediate percutaneous coronary intervention (PCI) in 155 interventional cardiology centers in Poland were included in this analysis. Patients with incomplete records concerning baseline characteristics were excluded. The reason for analyzing data from 2014 onwards was due to the fact that psoriasis was implemented for the first time into eCRF of the ORPKI database on January 1st, 2014.

Periprocedural mortality was defined as death from any cause during angiography/PCI. Psoriasis was defined as severe or moderate presentation as confirmed by prior diagnosis by a dermatologist (PASI index > 10). Only moderate and severe presentations were taken into account since they have been proven to have an impact on cardiovascular disease [6, 10, 12]. Allergic reaction was defined as any hypersensitive response of the immune system presenting as: bronchospasm, asthma exacerbation, conjunctivitis, urticaria,
eczema, angioedema, anaphylactic shock. All these symptoms had to be revealed during a patient stay at a catheterization laboratory and had not been present prior to admission.

Other complications or adverse events were diagnosed at the operator’s discretion according to current ESC definitions. Clinical outcome was defined as any periprocedural death. No further evaluation or follow-up of patients was performed. All patients provided informed consent. The study complied with ethical principles for clinical research based on the Declaration of Helsinki with later amendments. No external funding was used to support this registry.

**Statistical analysis**

Categorical variables were presented as numbers and percentages. The distributions of continuous variables were expressed as mean ± standard deviation (SD), standard error (SE), median with interquartile range (IQR). Additionally, the number of available cases and minimum and maximum values were also presented. Normality was assessed by the Lilliefors test. Equality of variances were assessed using the Levene test. Differences between groups were compared using the Student or Welch t-test depending on the equality of variances for normally distributed variables. The Mann-Whitney U test was used for non-normally distributed continuous variables. Categorical variables were compared with the Fisher exact test for the 2 × 2 tables or by the Pearson χ² test for the other tables. P-values less than 0.05 were assumed to indicate statistical significance.

Simple logistic regression was used for each of potential factor as a predictor for occurrence of allergic reaction. Then if p-value for a specific factor was less than 0.2 such factor was introduced into multiple logistic regression. The recommendation for using a significance level as high as 0.20 or 0.25 as screening criterion for initial variable selection is based on work by Bendel and Afifi (1977) on linear regression and on work by Mickey and Greenland (1989) on logistic regression. Only significant predictors and potential confounders (for adjustment) remained in the final model. The same procedure was repeated considering correction for rare events [18]. Both models were diagnosed using receiver operating characteristic curve and area under curve parameter.

The propensity score for psoriasis was calculated based on diabetes, hypertension, smoking, clinical diagnosis, kidney disease, presence of multivessel disease glycoprotein (GP) inhibitor use, access site, contrast volume, radiation, age, weight and gender. Then the odds
for allergic reaction were calculated using logistic regression with presence of psoriasis as a covariate and was conditioned on propensity score.


**Results**

There were 405,078 patients with full records in the ORPKI database for the period of January 1st 2014 till December 31st 2015 were included in this analysis. Psoriasis (moderate or severe) was diagnosed in 1507 (0.4%) of them. Patient baseline and clinical characteristics are presented in Table 1. Procedural outcome as well as frequency of complications in patients with and without psoriasis are presented in Table 2. The occurrence of periprocedural death, stroke and sudden cardiac arrest did not differ between the groups even though baseline and clinical characteristics in patients with psoriasis were high risk (significantly higher prevalence of diabetes, prior myocardial infarction, smoking and hypertension). Allergic reaction was more frequent during PCI in patients with psoriasis than in those without psoriasis (0.08% vs. 0.41%, p = 0.0226). There was no differences in the occurrence of periprocedural complications and death between the studied groups. Independent predictors of allergic reaction occurrence are presented in Table 3. Psoriasis was an independent predictor of allergic reaction occurrence (odds ratio [OR] 6.02; 95% confidence interval [CI] 1.44–25.22; p = 0.014). After propensity score adjustment psoriasis remained a significant predictor of allergic reaction (OR 5; 95% CI 1.2–20.7, p = 0.0245). Model adjustment for logistic regression is presented in Figure 1.

**Discussion**

Since psoriasis usually presents for the first time in adolescents and young adults, it is a lifelong burden and has a possible impact for development of other comorbidities. Psoriatic arthritis may appear in up to 30% of patients leading to decreased mobility and thus permanent disability which is an adverse condition for primary prevention of cardiovascular disease, metabolic syndrome and insulin resistance in diabetes mellitus [11]. The present study has confirmed more aggravating past medical history, comorbidities and higher risk demographics for patients with diagnosed psoriasis who were scheduled for coronary angiography and or PCI, either elective or urgent one. Mild psoriasis accounts for approx.
60% of all cases. The overall frequency of psoriasis (0.4%) was lower than in previous studies, however, the present focus was only on severe and moderate forms of psoriasis which have been documented to impact cardiovascular morbidity [10, 15, 19].

Recent coronary computed tomography studies revealed that patients with psoriasis had greater plaque burden and high risk plaque prevalence which was reduced at 1 year of observation following effective anti-inflammatory treatment which was revealing for the first time that such a correlation was observed [20]. Severe forms of psoriasis have also been named as a strong and independent predictor of ischemic stroke [21]. As previously observed, it seems that if any, psoriasis may have its deleterious effect only if it was at least expressed as moderate or severe presentation and this is the group which was under investigation herein. Even though patients with psoriasis in the registry were higher risk individuals with more frequent diabetes, hypertension and acute myocardial infarction (ST-segment elevation myocardial infarction [STEMI], non-ST-segment elevation myocardial infarction [NSTEMI]) presentation as well as multivessel and more pronounced atherosclerosis, their immediate periprocedural individual outcomes like death, stroke or sudden cardiac arrest were similar to those without psoriasis (< 1%). Presently it is believed that this may stem from the fact that a majority of complications related to psoriasis, if any, occur at a later stage and may be recorded only if a longer follow up period is available and only in severe forms of psoriasis [14, 15]. Adjusting for confounders, the presence of psoriasis does not seem to at least effect the periprocedural outcome of this group of patients.

The ORPKI database is one of the biggest datasets of consecutive patients with coronary artery disease treated invasively in European Union countries which continuously collects data on severe/moderate psoriasis occurrence since 2014. No previous studies with as large numbers of individuals as in the present registry (> 400,000 for a period of 2 consecutive calendar years) have been performed so far in investigating psoriasis as a comorbidity in the setting of percutaneous coronary interventions. Big data analysis in this case helped identify interesting associations such as more frequently observed allergic reaction occurrence in patients with psoriasis than in those without, which has remained an independent predictor even after statistical adjustments. It is known that several factors may contribute to anaphylaxis or hypersensitivity reactions during percutaneous coronary procedures, namely acetylsalicylic acid, P2Y12, heparin, GP IIb/IIIa inhibitors and contrast use itself [22–25]. Stent platforms and polymers used in DES stents may potentially induce such reactions as well [26, 27]. In the present study a majority of all these variables were
taken into consideration while performing multivariate analysis for independent predictors, however, not all were available in the database. The results indicate that GP IIb/IIIa inhibitor plays a significant role in allergic reaction occurrence as well as radiation doses which are a derivative of time of the procedure. It is worth noting that psoriasis conferred as the highest adjusted risk for allergic reaction even after propensity score adjustment. These findings need further investigation and suggest that caution should be used for patients presenting with severe or moderate forms of psoriasis when referred for invasive diagnostic and treatment. The results of the study may become hypotheses generating for future clinical studies. According to available research, this is the first report of such a relationship and requires further study.

**Limitations of the study**

The main limitation of this study is the non-randomized design. Only periprocedural outcome could be provided which limits the observation and incidence of complications that appeared in periods after the procedure. Statistical measures were undertaken to compensate for possible biases like propensity score matching and multivariate analysis in order to identify independent predictors but since this is an observational study not all possible confounders may have been identified. An underreporting of psoriasis may be possible but only moderate or severe forms (which are less frequent in general population) were gathered in ORPKI Registry. Finally, the diagnosis of psoriasis (moderate or severe) was based on prior recognition by a physician and no additional data concerning its treatment or systemic complications were gathered in this registry.

**Conclusions**

Severe or moderate psoriasis is a rare comorbidity in patients undergoing percutaneous coronary procedures (< 1%). However, it is an independent risk factor for allergic reaction occurrence during percutaneous coronary procedures. The adjusted periprocedural mortality and complications rates were similar in those with psoriasis versus those without.

**Conflict of interest:** None declared

**References**


<table>
<thead>
<tr>
<th>variable</th>
<th>No psoriasis</th>
<th>Psoriasis</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N = 403 571</td>
<td>N = 1507</td>
<td></td>
</tr>
<tr>
<td>Male gender</td>
<td>250,127 (62%)</td>
<td>1041 (69.1%)</td>
<td>&lt; 0.0001</td>
</tr>
<tr>
<td>Age</td>
<td>66.3 ± 10.9</td>
<td>63.3 ± 10.6</td>
<td>&lt; 0.0001</td>
</tr>
<tr>
<td>Diabetes</td>
<td>90,043 (22.3%)</td>
<td>492 (32.7%)</td>
<td>&lt; 0.0001</td>
</tr>
<tr>
<td>Prior MI</td>
<td>91,235 (22.6%)</td>
<td>383 (25.4%)</td>
<td>0.01</td>
</tr>
<tr>
<td>Prior PCI</td>
<td>102,131 (25.3%)</td>
<td>389 (25.8%)</td>
<td>0.656</td>
</tr>
<tr>
<td>Prior CABG</td>
<td>23,406 (5.8%)</td>
<td>81 (5.4%)</td>
<td>0.508</td>
</tr>
<tr>
<td>Smoking — active</td>
<td>68,486 (17%)</td>
<td>427 (28.3%)</td>
<td>&lt; 0.0001</td>
</tr>
<tr>
<td>Variable</td>
<td>No psoriasis</td>
<td>Psoriasis</td>
<td>( p )</td>
</tr>
<tr>
<td>---------------------------------------</td>
<td>--------------</td>
<td>-----------</td>
<td>--------</td>
</tr>
<tr>
<td>Hypertension</td>
<td>284,271 (70.4%)</td>
<td>1129 (74.9%)</td>
<td>0.0001</td>
</tr>
<tr>
<td>Indication:</td>
<td></td>
<td></td>
<td>0.0002</td>
</tr>
<tr>
<td>STEMI</td>
<td>48317 (11.9%)</td>
<td>219 (14.5%)</td>
<td></td>
</tr>
<tr>
<td>NSTEMI</td>
<td>52864 (13.1%)</td>
<td>240 (15.9%)</td>
<td></td>
</tr>
<tr>
<td>UA</td>
<td>124322 (30.8%)</td>
<td>440 (29.2%)</td>
<td></td>
</tr>
<tr>
<td>Stable angina</td>
<td>158635 (39.3%)</td>
<td>542 (35.9%)</td>
<td></td>
</tr>
<tr>
<td>Angiography only</td>
<td>221,327 (54.8%)</td>
<td>767 (50.9%)</td>
<td>0.0022</td>
</tr>
<tr>
<td>Multivessel disease with or without LMCA involvement</td>
<td>151,538 (37.7%)</td>
<td>643 (42.8%)</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>P2Y12 during the procedure</td>
<td>88,757 (21.9%)</td>
<td>397 (26.3%)</td>
<td>0.0002</td>
</tr>
<tr>
<td>UFH during PCI</td>
<td>153985 (84.5%)</td>
<td>644 (87%)</td>
<td>0.0595</td>
</tr>
<tr>
<td>LMWH during PCI</td>
<td>5980 (3.3%)</td>
<td>25 (3.4%)</td>
<td>0.8361</td>
</tr>
<tr>
<td>GP IIb/IIIa during PCI</td>
<td>12,056 (6.6%)</td>
<td>68 (9.2%)</td>
<td>0.0075</td>
</tr>
<tr>
<td>Thrombolysis during PCI</td>
<td>376 (0.2%)</td>
<td>1 (0.1%)</td>
<td>1.000</td>
</tr>
<tr>
<td>Contrast volume [mL]</td>
<td>128 ± 79</td>
<td>135 ± 83</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Radiation dose [Gy]</td>
<td>0.796 ± 0.952</td>
<td>0.957 ± 1.227</td>
<td>&lt;0.0001</td>
</tr>
</tbody>
</table>

CABG — coronary artery bypass grafting surgery; GP — glycoprotein; Gy — Grey; LMCA — left main coronary artery; LMWH — low molecular weight heparin; MI — myocardial infarction; NSTEMI — non-ST-segment elevation myocardial infarction; PCI — percutaneous coronary intervention; STEMI — ST-segment elevation myocardial infarction; UA — unstable angina; UFH — unfractionated heparin

**Table 2.** Clinical outcome and procedural complications.
Death during angiography 172 (0.08%)  0 (0%)  1.000
Death during PCI  983 (0.54%)  7 (0.95%)  0.128
SCA during angiography  905 (0.22%)  7 (0.46%)  0.0901
SCA during PCI  884 (0.49%)  6 (0.81%)  0.1818
Allergic reaction during angiography  160 (0.07%)  0 (0%)  1.000
Allergic reaction during PCI  145 (0.08%)  3 (0.41%)  0.0226
Stroke during angiography or PCI  55 (0.01%)  0 (0%)  1.000
Coronary artery perforation during PCI  295 (0.16%)  0 (0%)  0.6373
MI during PCI  173 (0.09%)  0 (0%)  1.000
No-reflow during PCI  1062 (0.58%)  8 (1.08%)  0.0850

MI — myocardial infarction; PCI — percutaneous coronary intervention; SCA — sudden cardiac arrest

**Table 3.** Independent predictors of allergic reaction during angiography and/or percutaneous coronary intervention

<table>
<thead>
<tr>
<th>Variable</th>
<th>OR</th>
<th>95% CI</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Psoriasis</td>
<td>6.02</td>
<td>1.44–25.22</td>
<td>0.014</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>0.47</td>
<td>0.24–0.92</td>
<td>0.026</td>
</tr>
<tr>
<td>Hypertension</td>
<td>0.57</td>
<td>0.36–0.89</td>
<td>0.014</td>
</tr>
<tr>
<td>STEMI/NSTEMI vs. other</td>
<td>0.58</td>
<td>0.35–0.97</td>
<td>0.036</td>
</tr>
<tr>
<td>GP IIb/IIIa inhibitors use</td>
<td>4.56</td>
<td>2.74–7.59</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>loge per 1 Gy (radiation dose)</td>
<td>1.67</td>
<td>1.22–2.28</td>
<td>0.001</td>
</tr>
<tr>
<td>Male gender</td>
<td>0.54</td>
<td>0.33–0.87</td>
<td>0.012</td>
</tr>
</tbody>
</table>
CI — confidence interval; GP — glycoprotein; Gy — Grey; STEMI — ST-segment elevation myocardial infarction; NSTEMI — non-ST-segment elevation myocardial infarction; OR — odds ratio

**Figure 1.** Model adjustment for logistic regression; *AUC — area under curve.