Disease Activity and Anti-VEGF Treatment Patterns in a Commercially Insured US Patient Population With Neovascular Age-Related Macular Degeneration

September 21, 2020

Arghavan Almony, MD¹ Katelyn R. Keyloun, PharmD, MS² Bijal Shah-Manek, PhD, B.Pharm³ Chi-Chang Chen, PhD, MsPharm⁴ Jasjit K. Multani, MPH⁴ Catherine B. McGuiness, MA, MSc⁴ Joanna Campbell, PhD²

¹Carolina Eye Associates, Southern Pines, NC, USA; ²Allergan, an AbbVie company, Irvine, CA, USA; ³Noesis Healthcare Technologies, Redwood City, CA, USA; ⁴IQVIA, Plymouth Meeting, PA, USA

Arghavan Almony, MD

- Consulting fee: Allergan, an AbbVie company
- Speakers Bureau: Allergan, an AbbVie company

Bijal Shah-Manek, PhD, BPharm

- Employee: Neosis Healthcare Technologies
- Consulting fee: Allergan, an AbbVie company, and Genentech self; Beigene, Cytokinetics, and Mirati spouse

Chi-Chang Chen, PhD, MsPharm, Jasjit K. Multani, MPH, Catherine B. McGuiness, MA, MSc

Employees: IQVIA, formerly QuintilesIMS

Katelyn R Keyloun, PharmD, MS; Joanna Campbell, PhD

• Employees: AbbVie Inc.

This study was sponsored by Allergan plc, Dublin Ireland (prior to its acquisition by AbbVie Inc.). Editorial assistance was provided to the authors by Michele Jacob, PhD, CMPP, of Evidence Scientific Solutions, Inc (Philadelphia, PA) and funded by AbbVie Inc. ICMJE authorship criteria were met. Neither honoraria nor payments were made for authorship.

Key Findings

- Among n=570 eyes presenting with active choroidal neovascularization (aCNV), 19.8% (n=113) transitioned to inactive CNV (iCNV) with a median transition time of 6.6 months
- Patients with aCNV incurred at least \$7,000 higher all-cause annual costs than patients with iCNV, largely driven by higher outpatient/anti-VEGF costs

Conclusions

- Rates of anti-VEGF treatment in this population were low compared with clinical trials, which
 may have contributed to limited transition to inactive/quiescent status
- Suboptimal treatment may lead to worse clinical outcomes (eg, worse visual acuity) and higher downstream costs
- Long-acting anti-VEGF therapy may help reduce treatment burden and preserve visual acuity

Limitation

 A key study limitation was the potential for miscoding in the claims date, which may have contributed to findings or lack of documentation of transitions of care

Background

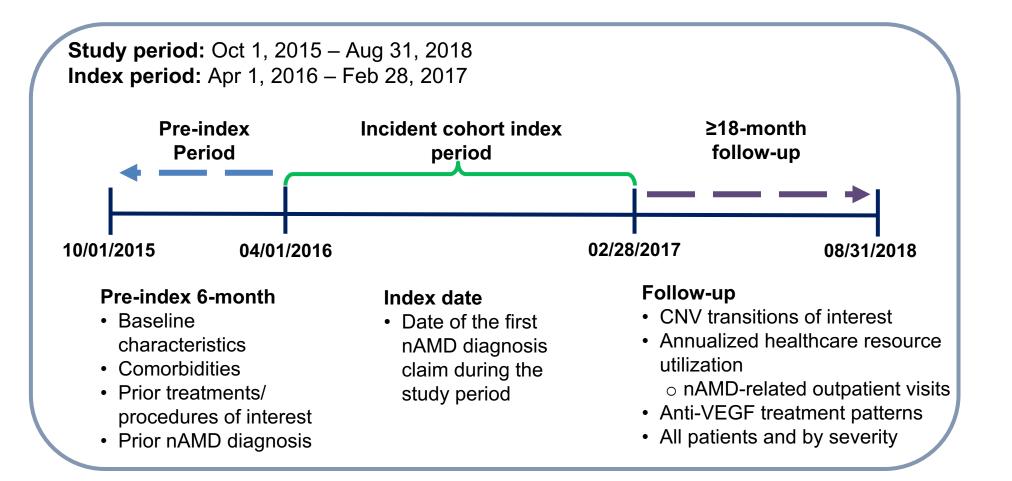
- Neovascular age-related macular degeneration (nAMD) accounts for ≤20% of AMD cases, but is responsible for ~90% of all cases of severe vision loss from the disease^{1,2}
- Available anti-VEGF therapies require frequent monitoring and regular intravitreal injections for optimal outcomes, resulting in a high patient, caregiver and healthcare system burden^{3,4}
- nAMD treatment patterns and transitions in disease status are not well understood in commercially insured US patients

Objective

 To assess choroidal neovascularization (CNV) activity and anti-VEGF treatment patterns among incident patients with nAMD in US clinical practice

Study Design

- Retrospective analysis of administrative claims data from IQVIA's PharMetrics® Plus database
 - Incident cohort: ≥50 years of age with commercial insurance, ≥1 claim(s) of ICD-10-CM nAMD diagnosis in the index period (per the figure), and ≥18 months of follow-up
 - Patients were stratified by disease status at diagnosis based on ICD-10-CM codes; ie, active CNV (aCNV), inactive/quiescent CNV (iCNV), inactive scar, and unspecified



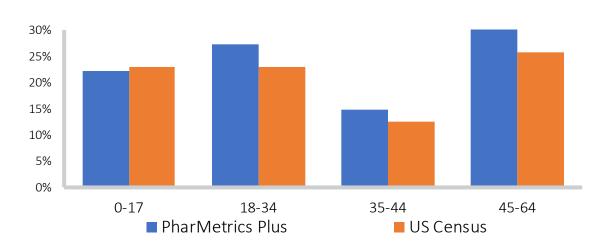
Outcome Measures

- Kaplan-Meier analyses were used to measure the time to first transition of disease status
 - From aCNV to iCNV
 - From iCNV to aCNV
 - From iCNV to inactive scar
- Mean annual healthcare resource utilization (HRU) is reported per patient
- Anti-VEGF treatment patterns were analyzed in the subgroup of patients who received ≥1 anti-VEGF treatment and had ≥12 months of follow-up after the initial treatment

PharMetrics® Plus Database

- One of the largest claims databases in the US with >150 million unique enrollees across all 50 states
- Representative of the national, commercially insured population in age and gender for people aged ≤65 years

PharMetrics Plus vs. US Census (Population Age)

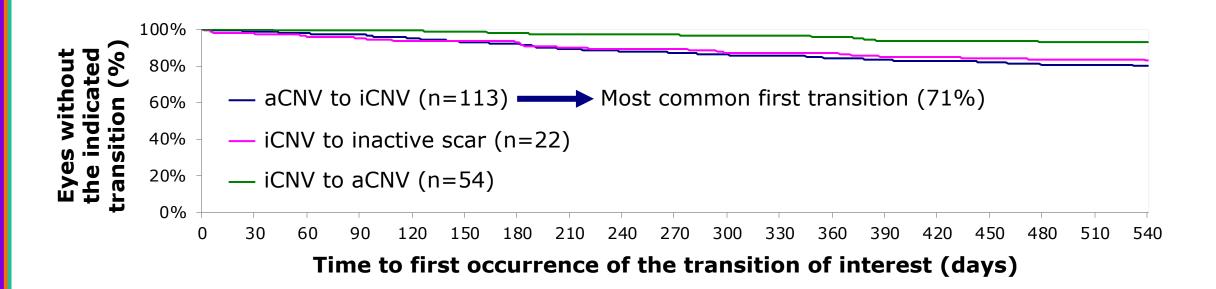


Of 1081 Incident nAMD Patients, Most had aCNV at Baseline and Few had Received Prior anti-VEGF Therapy

Parameter	aCNV n=501 (46.3%)	iCNV n=251 (23.2%)	Inactive scar n=124 (11.5%)	Unspecified n=205 (19.0%)
Mean (SD) age, y	67.3 (9.8)	66.8 (10.2)	70.7 (11.3)	67.2 (10.9)
Female, n (%)	280 (55.9)	145 (57.8)	62 (50.0)	111 (54.1)
Prior anti-VEGF use,a n (%)	13 (2.6)	1 (0.4)	0	8 (3.9)
Bevacizumab	0	0	0	0
Ranibizumab	4 (0.8)	0	0	3 (1.5)
Aflibercept	10 (2.0)	1 (0.4)	0	6 (2.9)
Prior fall/fracture, n (%)b	21 (4.2)	15 (6.0)	10 (8.1)	6 (2.9)
Mean (SD) CCI score	1.1 (1.5)	1.1 (1.8)	1.3 (1.7)	1.3 (1.6)
Ocular comorbidities, n (%)				
DR	29 (5.8)	7 (2.8)	6 (4.8)	14 (6.8)
OAG/OHT	28 (5.6)	19 (7.6)	14 (11.3)	15 (7.3)
OSD	55 (11.0)	11 (4.4)	7 (5.6)	13 (6.3)
Uveitis	9 (1.8)	2 (0.8)	1 (0.8)	0

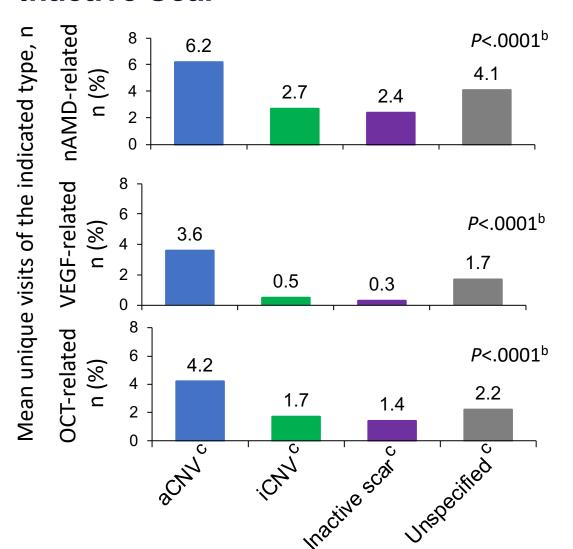
^a Not mutually exclusive. ^b During the 6-month pre-index period CCI, Charlson Comorbidity Index; DR, diabetic retinopathy; OAG, open-angle glaucoma; OHT, ocular hypertension; OSD, ocular surface disease

Most aCNV Eyes Remained Without a Transition Through 18 Months



- The most common baseline status was aCNV (45%, N=570) of 1270 incident eyes diagnosed with nAMD at baseline
- Among eyes with active CNV at baseline, only 28% (n=160) had any transition and only 20% transitioned to inactive CNV (n=113), the most common first transition
- The median time in eyes transitioning from active to inactive CNV was \sim 6.6 months (mean time of \sim 7.6 months)

Annual nAMD-Related Healthcare Resource Utilization and Costs Were Significantly Higher in Patients With aCNV Than Those With iCNV or Inactive Scar^a



Direct Costs (per patient)*	Active CNV (n= 501) Annual Cost, Mean (SD)	Inactive CNV (n=251) Annual Cost, Mean (SD)
Total annual All-cause cost	\$21, 352 (\$33,040)	\$14,030 (\$22,399)
All-cause Outpatient cost	\$13,103 (\$15,428)	\$7,241 (\$11,106)
 nAMD related outpatient Anti-VEGF related outpatient OCT -related outpatient costs 	\$5,287 (\$8,028)\$3,894 (\$6,870)\$247 (\$ 252)	\$985 (\$2,613)\$471 (\$2,261)\$96 (\$103)
All-cause Pharmacy cost	\$ 3,611 (\$7,917)	\$2,787 (\$4,563)
All-cause Hospitalization costs	\$4,087 (\$22,729)	\$3,701 (\$14,960)
All-cause ER costs	\$552 (\$2,114)	\$301 (\$958)

Patients with active choroidal neovascularization (CNV) incurred at least \$7,000 higher all-cause annual costs than patients with inactive CNV, largely driven by higher outpatient/anti-VEGF costs

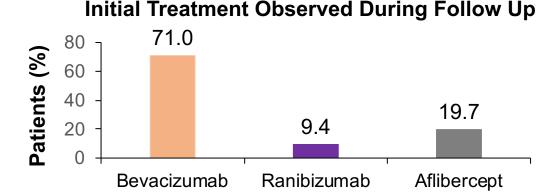
^a Data reported per patient. ^b Across severity subgroups. Statistical analysis included the "Unspecified" disease subgroup. ^c Total, N=1081 (100%); aCNV, 501 (46.3%); iCNV, n=251 (23.2%); Inactive scar, n=124 (11.5%); Unspecified, n=205 (19.0%). OCT, optical coherence tomography.

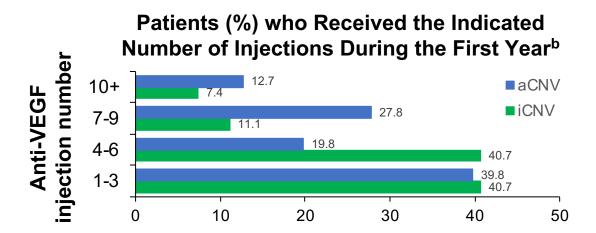
Anti-VEGF Treatment Patterns

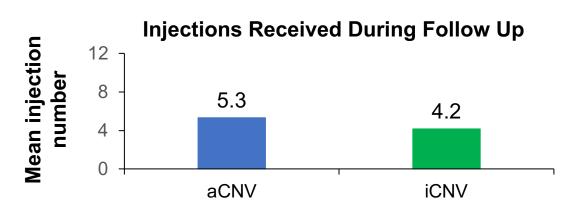
 427 incident patients receiving anti-VEGF treatment were included in this subgroup analysis^a

- ~41% of patients received 1-3 anti-VEGF treatments during follow up
 - Overall, ~62% of patients received ≤6 injections over the first year

 The overall mean (SD) number of injections during one year of follow up was 5.2 (3.5)



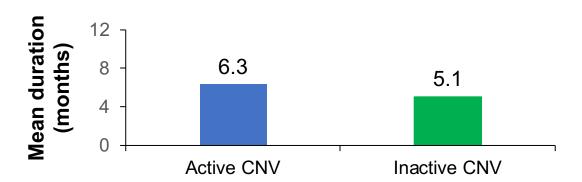




^a Patients treated with only 1 anti-VEGF type on initial injection date and during 12-months post-initial injection. ^b Including initial injection

Anti-VEGF Treatment Patterns Analysis (continued)

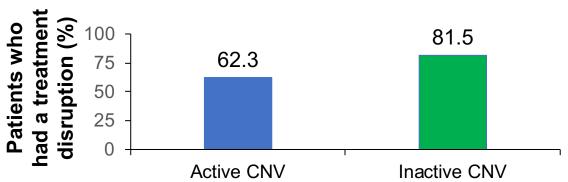
 Overall, the mean (SD) duration of therapy was 6.2 (4.7) months^a



 Overall, 282 (66.0%) patients reported a treatment disruption/ break^b

Treatment Received During Follow Up

Treatment Received During Follow up



^a Time from diagnosis to the last recorded anti-VEGF injection.

^b Defined as a >18-week gap in any anti-VEGF therapy.

Key Findings

- Among n=570 eyes presenting with active choroidal neovascularization (aCNV), 19.8% (n=113) transitioned to inactive CNV (iCNV) with a median transition time of 6.6 months
- Patients with aCNV incurred at least \$7,000 higher all-cause annual costs than patients with iCNV, largely driven by higher outpatient/anti-VEGF costs

Conclusions

- Rates of anti-VEGF treatment in this population were low compared with clinical trials, which
 may have contributed to limited transition to inactive/quiescent status
- Suboptimal treatment may lead to worse clinical outcomes (eg, worse visual acuity) and higher downstream costs
- Long-acting anti-VEGF therapy may help reduce treatment burden and preserve visual acuity

Limitation

 A key study limitation was the potential for miscoding in the claims date, which may have contributed to findings or lack of documentation of transitions of care