

Pearls for Anti-Coagulation Optimization in Older Adults

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Learning Objectives

1. Describe common indications for anticoagulation therapy in older adults in PALTC

2.Identify risks and benefits of anticoagulation therapy in older adults in PALTC

3. Criticize approaches to different anticoagulation treatments in PALTC



Disclosures

None

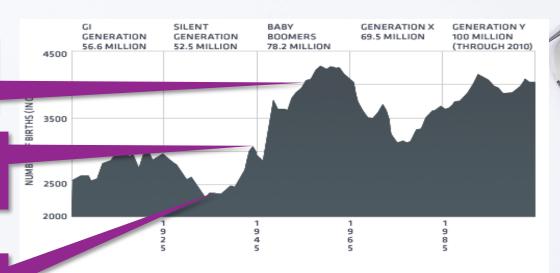


US BIRTHS 1905-2002



2022 Born 1942 80 yoa

2018 Born 1938 80 yoa



GENERATION: 1905-1924 56.6 MILLION
SILENT GENERATION: 1925-1944 52.5 MILLION
BABY BOOMERS: 1945-1964 78.2 MILLION
GENERATION X: 1965-1984 69.5 MILLION
GENERATION Y: 1985-2004 79.5 MILLION
GENERATION Z: 2005-2024 = 16 MILLION (THUSFAR)
GENERATION BLEND*: 2025-2044! UNKNOWN

*ACC HAS CHOSED THE MONIKER "GENERATION BLEND" BECAUSE THIS WILL BE THE MOST ETHNICALLY ASSIMILATED U.S. GENERATION EVER.



Anticoagulation is common in long term care.

87% of strokes are ischemic¹

- Atrial Fibrillation impacts
 - 5.2 million today²
 - 12.1 million people by 2030³
- Treatment on Venous Thromboembolism (VTE)
- Prophylaxis of VTE



^{3.} Colilla S, Crow A, Petkun W, Singer DE, Simon T, Liu X. Estimates of current and future incidence and prevalence of atrial fibrillation in the U.S. adult population. *Am J Cardiol.* 2013;112:1142–1147. doi: 10.1016/j.amjcard.2013.05.063.



Anticoagulation is common in long term care.

A large study in 2010 evaluated warfarin use for Afib in LTC¹

- INRs were suboptimal
- 17%-57% of Afib residents on Warfarin
- Challenges outweigh benefits

Rates in a study in LTC in 2017²

- 38% Warfarin
- 32% NOACs
- 30% None



^{1.} Neidecker M, Patel AA, Nelson WW, Reardon G. Use of warfarin in long-term care: a systematic review. *BMC Geriatr*. 2012;12:14. Published 2012 Apr 5. doi:10.1186/1471-2318-12-14

^{2.} Rojas-Fernandez CH, Goh J, Hartwick J, Auber R, Zarrin A, Warkentin M, Hudani Z. Assessment of Oral Anticoagulant Use in Residents of Long-Term Care Homes: Evidence for Contemporary Suboptimal Use. Ann Pharmacother. 2017 Dec;51(12):1053-1062. doi: 10.1177/1060028017723348. Epub 2017 Jul 26. PMID: 28745065.

"I Walk the Line"



Warfarin is the most common drug involved in error in 2012.1

- 32,176 medication error incidents reported over a 2-year period
 - 1,623 (5%) were anticoagulant medication errors
 - 2% of these errors (n=29) resulted in patient harm.
 - Anticoagulant medication errors had higher odds of patient harm when compared with other errors (OR 1.79)

The most litigious medication in long term care

- The nursing staff does not ensure the labs are drawn daily as ordered.
- The physician forgets to order the daily labs.
- The nurse does not report the labs to the physician and the two medications are used together for too long.



1. Desai, R., Williams, C.E., Greene S.B., Pierson S. and Hansen R.A. "Anticoagulant medication errors in nursing homes: characters, causes, outcomes and association with patient harm", Journal of Healthcare Risk Management, (accepted for publication November 2012)

Use of NOACs in LTC

Effectiveness and safety of oral anticoagulants in elderly patients with atrial fibrillation

- 30 401 patients ≥75 years identified (median age 82 years, 53% women, mean CHA₂DS₂-VaSc score 4.5)
- Reduced (renal) doses in 49% of patients
- Efficacy similar between NOACs and warfarin
- Bleeding risk reduced or similar with NOACs vs. warfarin



Rutherford OW, Jonasson C, Ghanima W, et al Effectiveness and safety of oral anticoagulants in elderly patients with atrial fibrillation *Heart* 2022;**108**:345-352.

Comparison of NOACs



RE-LY (dabigatran)	ROCKET-AF (rivaroxaban)	ARISTOTLE (apixaban)	ENGAGE AF-TIMI 48 (edoxaban)
18,113	14,264	18,201	21,105
Patients with NVAF CHADS₂ score ≥1 (mean 2.1) Mean age: 72 years	Patients with NVAF CHADS₂ score ≥2 (mean 3.5) Mean age: 73 years	Patients with NVAF CHADS₂ score ≥1 (mean 2.1) Mean age: 70 years	Patients with NVAF CHADS₂ score ≥2 (mean 2.8) Mean age: 72 years
Double-blind randomized, non-inferiority trial	Double-blind randomized, non-inferiority trial	Double-blind randomized, non-inferiority trial	Double-blind randomized, non-inferiority trial
150 mg (110 mg) twice daily	20 mg (15 mg) once daily	5 mg (2.5 mg) twice daily	60 mg (30 mg) once daily
Warfarin (INR 2-3) TTR 64%	Warfarin (INR 2-3) TTR 55%	Warfarin (INR 2–3) TTR 62%	Warfarin (INR 2-3) TTR 68.4%
Stroke (ischemic or hemorrhagic) or systemic embolism	Stroke (ischemic or hemorrhagic) or systemic embolism	Stroke (ischemic or hemorrhagic) or systemic embolism	Stroke (ischemic or hemorrhagic) or systemic embolism
Major bleeding	Composite of major and non-major bleeding	Major bleeding	Major bleeding
Efficacy of dabigatran 110 mg vs. warfarin (0.91; 95% CI, 0.74–1.11; P<0.001 for non-inferiority) Efficacy of dabigatran 150 mg vs. warfarin (0.66; 95% CI, 0.53–0.82; P<0.001 for superiority) Safety of dabigatran 110 mg vs. warfarin (0.80; 95% CI,	Efficacy of rivaroxaban 20 mg vs. warfarin (0.88; 95% Cl, 0.74–1.03; P<0.001 for non-inferiority; P=0.12 for superiority) Safety of rivaroxaban 20 mg vs. warfarin (1.03; 95% Cl, 0.96–1.11; P=0.44)	Efficacy of apixaban 5 mg vs. warfarin (0.79; 95% CI, 0.66–0.95; P<0.001 for non-inferiority; P=0.01 for superiority) Safety of apixaban 20 mg vs. warfarin (0.69; 95% CI, 0.60–0.80; P<0.001)	Efficacy of edoxaban 60 mg vs. warfarin (0.87; 97.5% CI, 0.73–1.04; P=0.08 for superiority Efficacy of edoxaban 30 mg vs. warfarin (1.13; 97.5% CI, 0.96–1.34; P=0.10 for superiority Safety of edoxaban 60 mg vs warfarin (0.80; 95% CI,
0.69–0.93; P=0.003) Safety of dabigatran 150 mg vs. warfarin (0.93; 95% CI,			0.71–0.91; P<0.001) Safety of edoxaban 30 mg vs warfarin (0.47; 95% CI, 0.41–0.55; P<0.001)
	(dabigatran) 18,113 Patients with NVAF CHADS₂ score ≥1 (mean 2.1) Mean age: 72 years Double-blind randomized, non-inferiority trial 150 mg (110 mg) twice daily Warfarin (INR 2–3) TTR 64% Stroke (ischemic or hemorrhagic) or systemic embolism Major bleeding Efficacy of dabigatran 110 mg vs. warfarin (0.91; 95% CI, 0.74–1.11; P<0.001 for non-inferiority) Efficacy of dabigatran 150 mg vs. warfarin (0.66; 95% CI, 0.53–0.82; P<0.001 for superiority) Safety of dabigatran 110 mg vs. warfarin (0.80; 95% CI, 0.69–0.93; P=0.003) Safety of dabigatran 150 mg	(dabigatran) (rivaroxaban) 18,113 14,264 Patients with NVAF CHADS₂ score ≥1 (mean 2.1) Patients with NVAF CHADS₂ score ≥2 (mean 3.5) Mean age: 72 years Mean age: 73 years Double-blind randomized, non-inferiority trial Double-blind randomized, non-inferiority trial 150 mg (110 mg) twice daily Warfarin (INR 2-3) TTR 64% Stroke (ischemic or hemorrhagic) or systemic embolism Stroke (ischemic or hemorrhagic) or systemic embolism Major bleeding Stroke (ischemic or hemorrhagic) or systemic embolism Efficacy of dabigatran 110 mg vs. warfarin (0.91; 95% CI, 0.74-1.11; P<0.001 for non-inferiority) Composite of major and non-major bleeding Efficacy of ivaroxaban 20 mg vs. warfarin (0.88; 95% CI, 0.74-1.03; P<0.001 for non-inferiority; P=0.12 for superiority) Safety of dabigatran 110 mg vs. warfarin (0.80; 95% CI, 0.69-0.93; P=0.003) Safety of dabigatran 150 mg Safety of dabigatran 150 mg (1.03; 95% CI, 0.96-1.11; P=0.44)	(dabigatran)(rivaroxaban)(apixaban)18,11314,26418,201Patients with NVAF CHADS₂ score ≥1 (mean 2.1)Patients with NVAF CHADS₂ score ≥2 (mean 3.5)Patients with NVAF CHADS₂ score ≥2 (mean 3.5)Double-blind randomized, non-inferiority trialDouble-blind randomized, non-inferiority trialDouble-blind randomized, non-inferiority trialDouble-blind randomized, non-inferiority trialDouble-blind randomized, non-inferiority trialDouble-blind randomized, non-inferiority trial150 mg (110 mg) twice daily20 mg (15 mg) once daily5 mg (2.5 mg) twice dailyWarfarin (INR 2-3) TTR 64%Warfarin (INR 2-3) TTR 55%TTR 62%Stroke (ischemic or hemorrhagic) or systemic embolismStroke (ischemic or hemorrhagic) or systemic embolismStroke (ischemic or hemorrhagic) or systemic embolismMajor bleedingComposite of major and non-inferiorityMajor bleedingEfficacy of dabigatran 110 mg vs. warfarin (0.91; 95% CI, 0.74-1.11; P<0.001 for non-inferiority)Efficacy of rivaroxaban 20 mg vs. warfarin (1.03; 95% CI, 0.96-1.11; P=0.44)Efficacy of apixaban 5 mg vs. warfarin (0.66; 95% CI, 0.96-0.93; P<0.001 10.80; 95% CI, 0.96-1.11; P=0.44)Safety of dabigatran 110 mg vs. warfarin (0.80; 95% CI, 0.69-0.93; P=0.003)Safety of rivaroxaban (1.03; 95% CI, 0.96-1.11; P=0.44)Efficacy of apixaban 20 mg vs. warfarin (0.69; 95% CI, 0.60-0.80; P<0.001)

ARISTOTLE, Apixaban for reduction in Stroke and Other Thromboembolic Events in Atrial Fibrillation; CI, confidence interval; ENGAGE AF-TIMI 48, Effective Anticoagulation with Factor Xa Next Generation in Atrial Fibrillation-Thrombolysis in Myocardial Infarction 48; NVAF, mon-valvular atrial fibrillation; RE-LY, Randomized Evaluation of Long-Term Anticoagulation Therapy trial; ROCKET-AF, Rivaroxaban Oncedaily, oral direct factor Xa inhibition Compared with vitamin K antagonism for prevention of stroke and Embolism Trial in Atrial Fibrillation; TTR, mean percent of time in the therapeutic range. Other abbreviations as in Table 3.

Indolfi, Ciro & Santarpia, Giuseppe & Curcio, Antonio & Sibilio, Gerolamo. (2015). Atrial Fibrillation and anticoagulation. 10.13140/RG.2.1.2372.5602.

Cost

- Warfarin \$20 a month
- Dabigatran \$475 a month (generic in June? 2022)
- Rivaroxaban \$550 a month (generic in litigation)
- Apixaban \$550 a month (generic after 2026)
- Edoxaban \$380 a month (Generic)



Covid-19



- For non-hospitalized patients with COVID-19, anticoagulants and antiplatelet therapy should not be initiated for prevention of venous thromboembolism (VTE) or arterial thrombosis unless there are other indications (AIII).
- Hospitalized adults with COVID-19 should receive VTE prophylaxis per the standard of care for other hospitalized adults (AIII).
- Hospitalized patients with COVID-19 should not routinely be discharged on VTE prophylaxis (AIII).
- Using Food and Drug Administration-approved regimens, extended VTE prophylaxis can be considered in patients who are at low risk for bleeding and high risk for VTE as per protocols for patients without COVID-19 (BI)

Q&A



