

Introduction

- Colorectal cancer (CRC) incidence is rising, particularly among those 20-49 years old.
- Approximately 10.5% of new CRC diagnoses occur in those under 50 years old.
- Individuals with early-onset CRC should be referred for cancer genetic testing to evaluate for an inherited syndrome.
- Genetic referral rates for this target population have been low.
- Hereditary CRC is commonly inherited in an autosomal dominant pattern although some syndromes (*MUTYH* associated polyposis and *NTHL1*) are inherited in an autosomal recessive pattern.

Methods

- Retrospective chart review of individuals age ≤ 50
- Cancer registry database with colon or rectal adenocarcinoma from 2016 to 2020.
- The primary intervention was to contact via telephone those who had not been previously referred for cancer genetic testing and to invite them to undergo risks assessment and testing.

Baseline Demographics

Variable	Previously Referred	Eligible for Referral
N	34	36
Average Age	45	43.5
Caucasian	23	19
African American	5	7
Non-White Hispanic	4	7
Asian	2	3

Results

Variable	Pre Intervention	Post Intervention	% Change	p-Value
N	34	44	29.4	0.787
Caucasian	23	29	26.1	0.178
African American	5	6	20.0	0.201
Non-White Hispanic	4	7	75	0.682
Asian	2	2	0	1.00

- Overall, referrals to cancer genetics increased 29.4%.
- Differences in referral rates across races exist.

Conclusion

- A subset of young onset CRC may be detected earlier by identifying those with associated mutations.
- Helping patients navigate using age criteria and tumor subtype may increase referral rates to genetics counselors.
- Racial disparities in genetics referrals exist, but improving navigation can increase referrals across all races.
- A detailed history of first and second degree relatives may identify at risk patients.

Future Directions

- Prospective best practice advisory to identify patients who qualify for referral.

References

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