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INTRODUCTION

Regorafenib for the treatment of patients with previously treated **metastatic colorectal cancer (mCRC)**

PURPOSE

To evaluate the **efficacy** and **safety** of **regorafenib** treatment in patients with **mCRC**

MATERIAL AND METHODS

- ✓ Retrospective observational **study**
- ✓ mCRC patients treated with **regorafenib** (September 2015-2017)
- ✓ **Collected variables**: age, sex, ECOG, KRAS gene status, treatment line, number of cycles, dose reduction
- ✓ **Efficacy endpoints**: progression-free survival (PFS) and overall survival (OS) obtained by the Kaplan-Meier method
- ✓ **Safety**: collected adverse effects (AE)
- ✓ **Descriptive statistical analysis**: SPSS®Statistics program V22.0

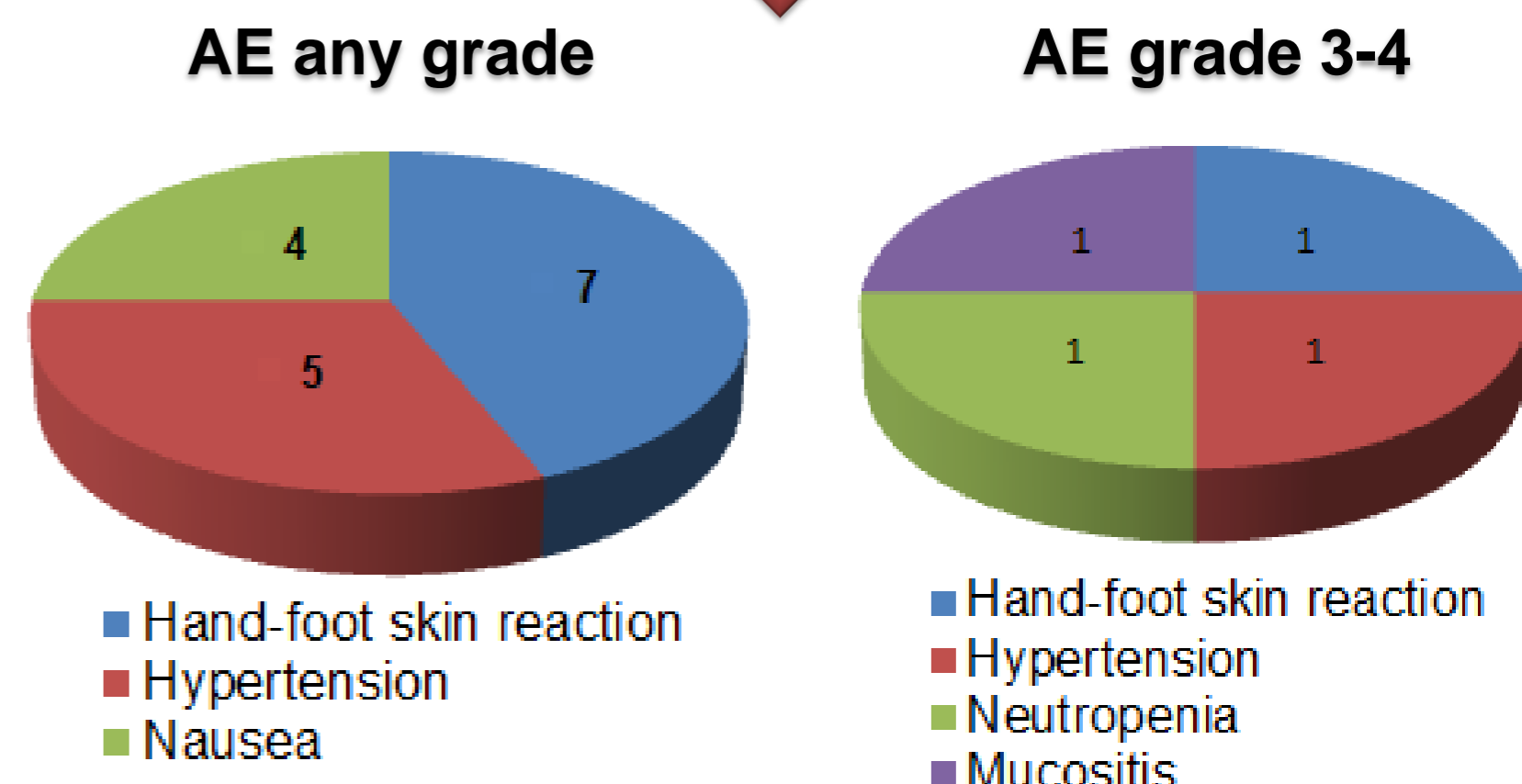
RESULTS

| | |
|----------------------------------|--|
| Patients | n=13 (7 men; 6 women) |
| Age | Median=57 years (41-77) |
| Initial ECOG | 0=38,46 % 1=38,46% 2=23% |
| KRAS gene | Mutated=50,8% Wild-type=30,8% Undetermined=3,8% |
| Protocol | Regorafenib 160 mg once daily for 21 days, every 28 days |
| Line | 3rd=46,2% >4th=53,8% |
| Dose reduction | 30,8% |
| Nº of cycles | Mean=2.75±1.22 cycles |
| Treatment discontinuation | n=9 progression n=3 deterioration of general health n=1 toxicity |

EFFICACY



SAFETY



CONCLUSION

- ✓ The **SLP** obtained is greater than what was described in the pivotal trial CORRECT (**3.0 versus 1.9 months**). This was possibly due to the longer time it took to determine the radiological response.
- ✓ The **SG** was also higher (**8.3 versus 6.4 months**) → limitation of the sample size.
- ✓ The **AE** described were similar to those published in the literature.