The use of recovery animals across monoclonal antibody development packages: opportunity for further optimization remains

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INTRODUCTION

- It is a regulatory requirement that recovery of adverse findings is assessed during pharmaceutical development, but there is flexibility around how and when this is performed and if recovery animals are necessary.
- For monoclonal antibodies (mAbs) following ICH S6(R1), if use of recovery animals is warranted, this need only be in one toxicity study.
- We have used data shared within a recent collaboration between the NC3Rs, the Netherlands Medicines Evaluation Board (MEB) and 14 pharmaceutical companies to review current practices for recovery animals use during mAb development.

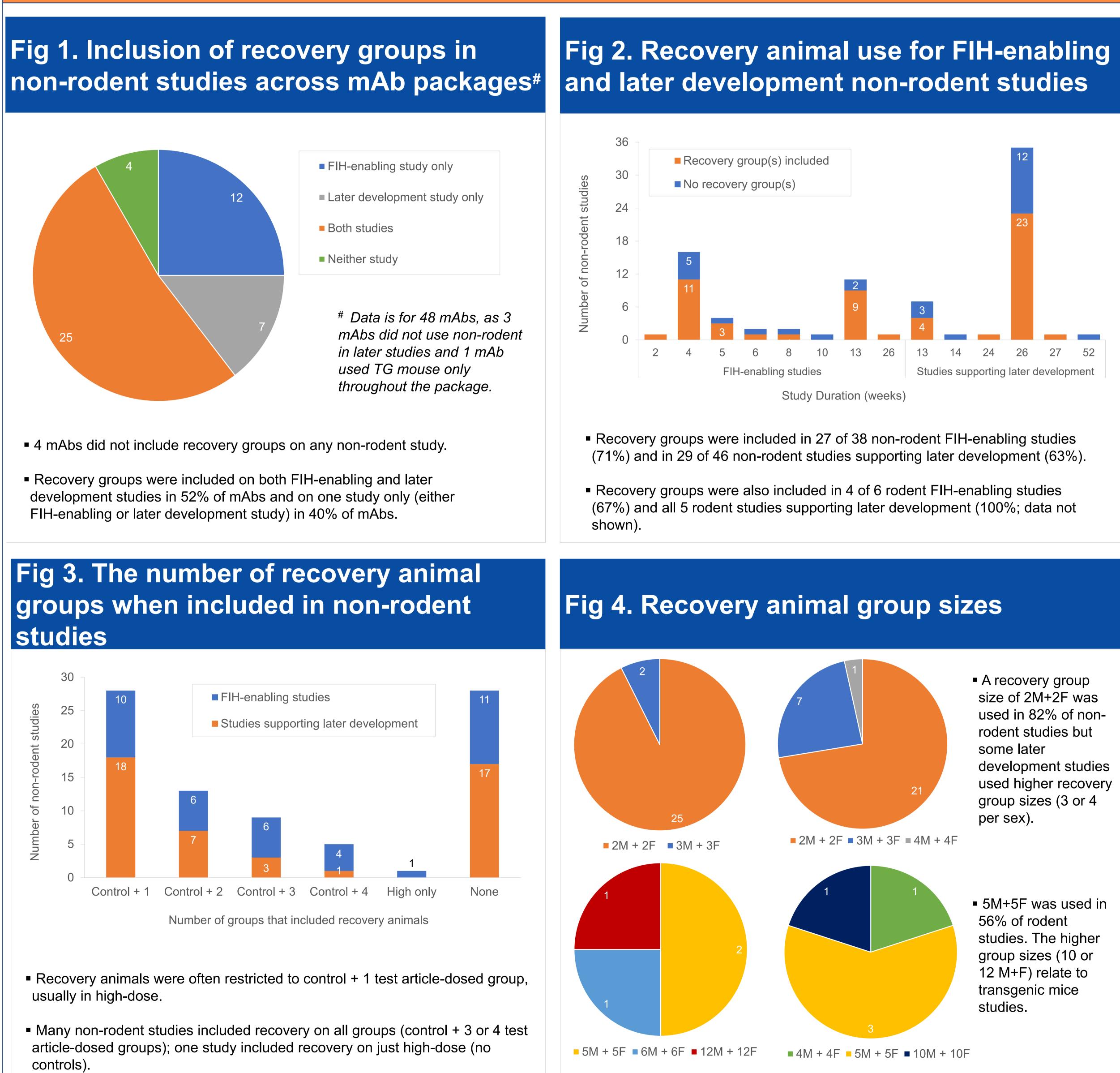
METHODS

- Data on study designs (e.g., start date, species, recovery animal group number and sizes), for studies enabling first-in-human (FIH) dosing and longer duration studies supporting later development were collected.
- To compare with previous data [1], only mAbs with at least one study started in 2015 or later were used in this analysis; there were 52 mAbs with 83 non-human primate, 1 minipig, 4 rat, 4 mouse and 3 transgenic (TG) mouse studies in total.

[1] Sewell F et al. (2014). Reg Tox & Pharm 70: 413-429.

CONCLUSION

- Variability in study designs suggests case-by-case approaches are used to develop many mAbs.
- Recovery is often assessed in multiple studies and multiple species.
- These data suggest assessment of recovery is more extensive than required by ICH S6(R1), and there may be an opportunity to reduce recovery animal use on many mAb programs.



studies



- not shown).

There was also one rodent study with recovery on low-dose group only (data)

Top panel: Non-rodent studies; Lower panel: rodent studies. FIH-enabling studies on left and Later development studies on right.

Abstract 4501; Poster P509

RESULTS

Table 1. Recovery animal use for the 8 mAbs using two species across the package

mAb ID	FIH-enabling studies	Later development studies
1	Rat (13 wk ✓) + Cynomolgus monkey (13 wk ✓)	Rat (26 wk ✓)
2	Mouse (13 wk ✓) + Cynomolgus monkey (13 wk ✓)	Cynomolgus monkey (26 wk X)
3	Mouse (2 wk X) + Cynomolgus monkey (4 wk X)	Mouse (13 wk ✓) + Cynomolgus monkey (13 wk ✓)
4	Rat (8 wk <mark>X</mark>) + Cynomolgus monkey (8 wk ✓)	Cynomolgus monkey (26 wk X)
5	Rat (8 wk ✓) + Minipig (6 wk ✓)	Rat (26 wk ✓)
6	TG mouse (5 wk <u>X</u>) + Cynomolgus monkey (5 wk <u>X</u>)	TG mouse (26 wk ✓) + Cynomolgus monkey (26 wk ✓)
7	Rat (4 wk ✓) + cynomolgus monkey (4 wk ✓)	Rat (26 wk ✓) + Cynomolgus monkey (26 wk ✓)
8	Mouse (13 wk ✓) + Cynomolgus monkey (13 wk ✓)	Mouse (26 wk ✓) + Cynomolgus monkey (26 wk ✓)

(X wk): study duration recovery groups included X no recovery groups

only).

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4 mAbs included recovery groups in all studies (both species), 3 mAbs included recovery groups in both species but for only one study duration and 1 mAb included recovery animals on only one study (non-rodent

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