

| | b vs. Placebo as add-on treatment for OPD patients characterized by eosinophil level |
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| Results information | |
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| Trial information | |
| Trial identification | |
| Additional study identifiers | |
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| Sponsors | |
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| Paediatric regulatory details | |
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| Results analysis stage | | | | |
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| Subject disposition | | | | |
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| Number of subjects in period 1[1] | | |
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Baseline characteristics Reporting groups Reporting group values Reporting group values

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Secondary: Rate of COPD exacerbations requiring emergency department (ED) visits

| and/or hospitalizations | | | | |
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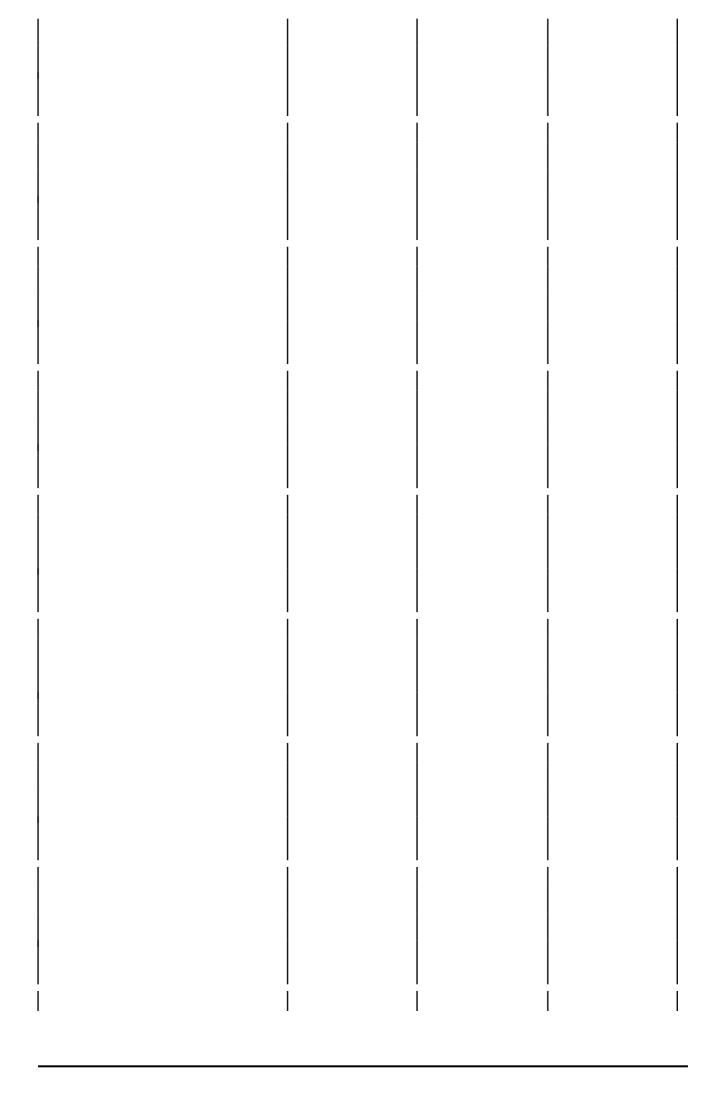
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| Socondary Change from Pagelin | o in Maar CO | DD accoccina | nt tost (CAT) | ccorc |
| Secondary: Change from Baseline | e in Mean CO | מאפטניטוופו | it test (CAT) | |
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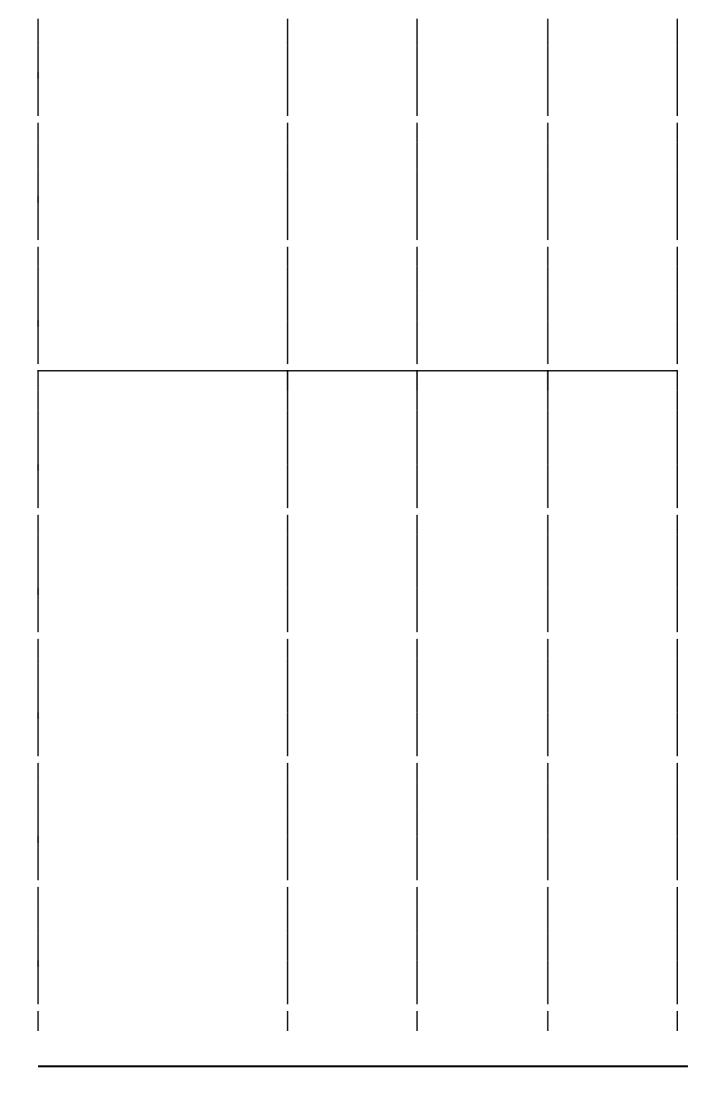
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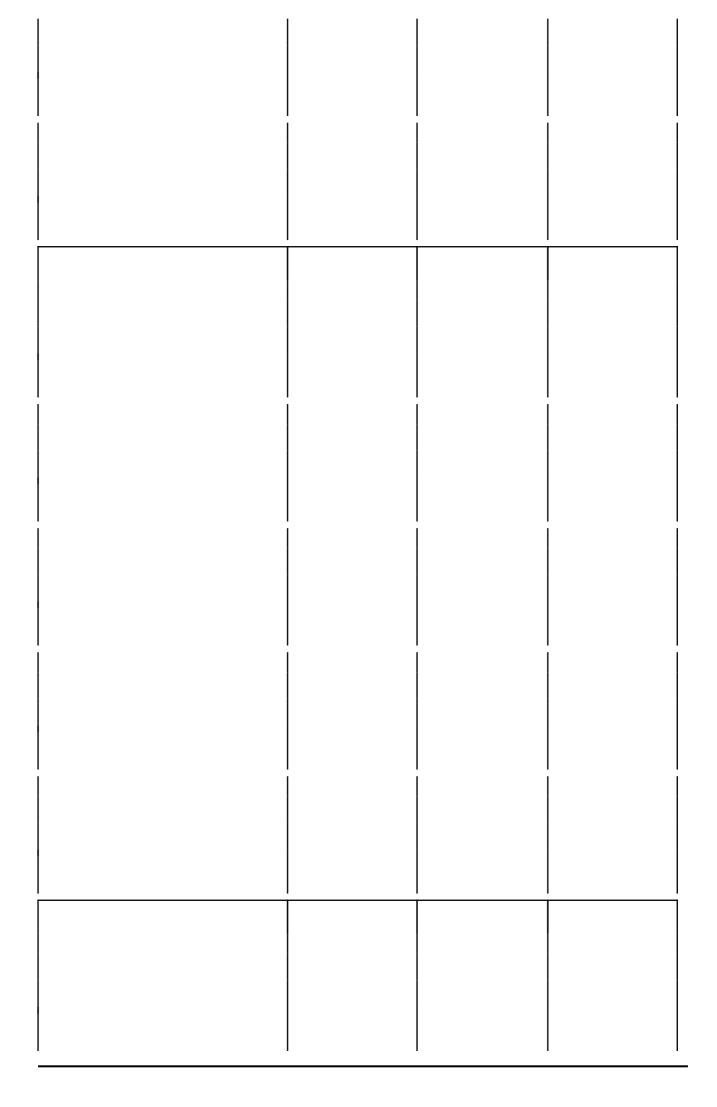
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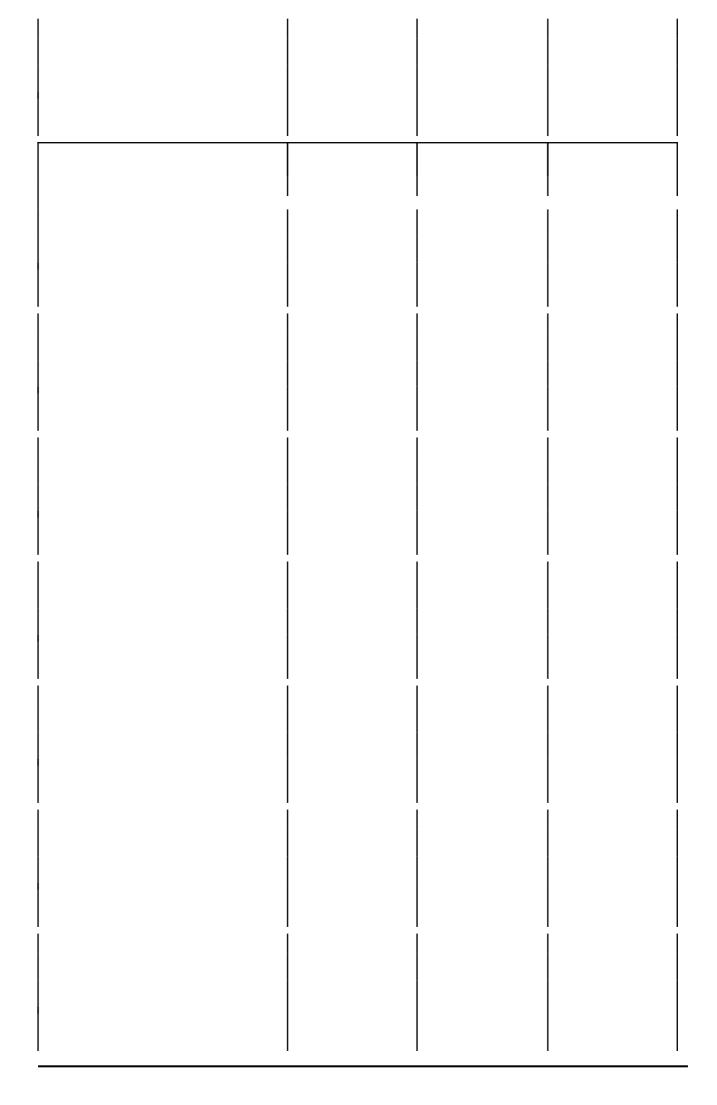
Adverse events information Dictionary used Reporting groups Serious adverse events

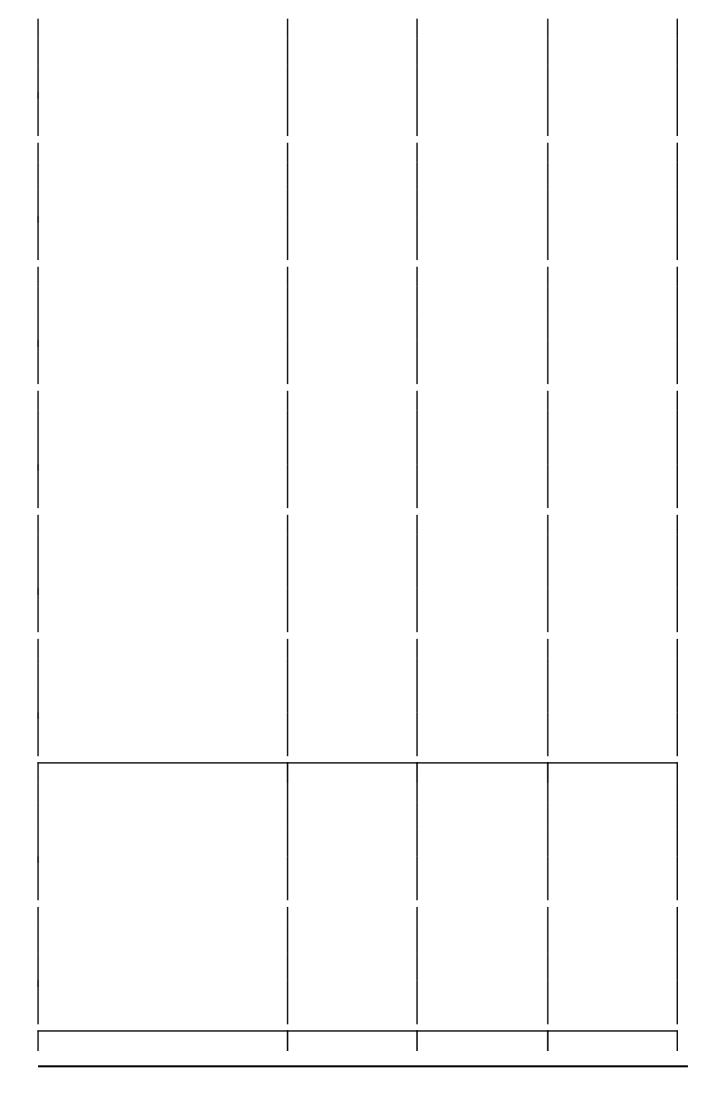
Adverse events

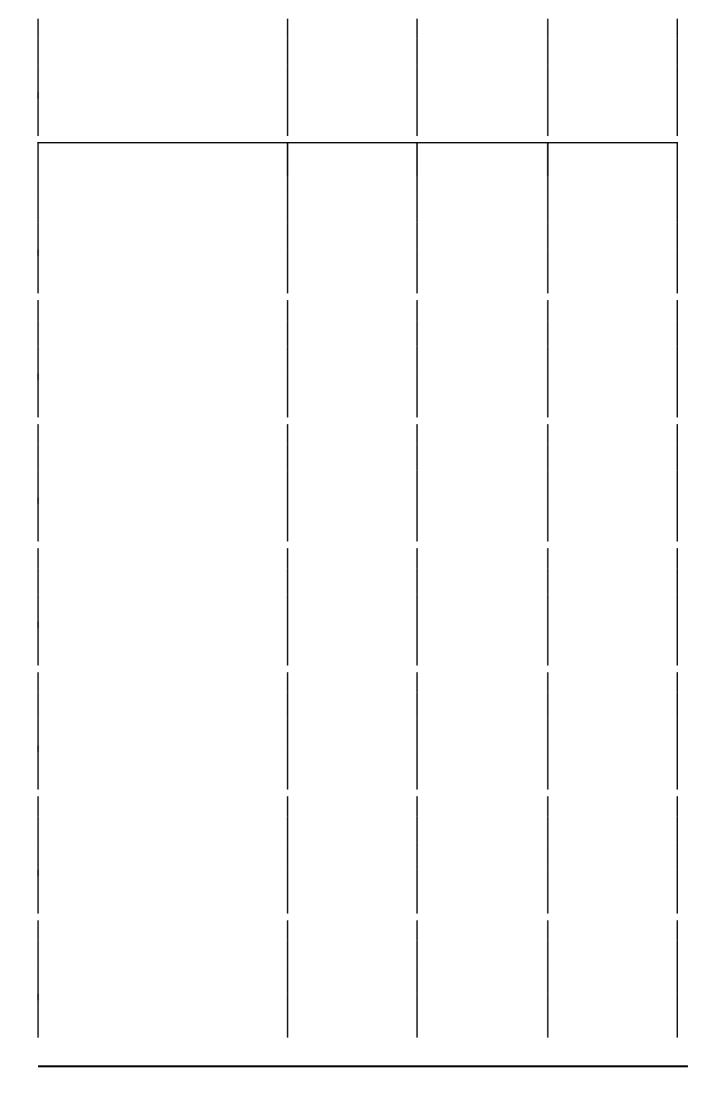


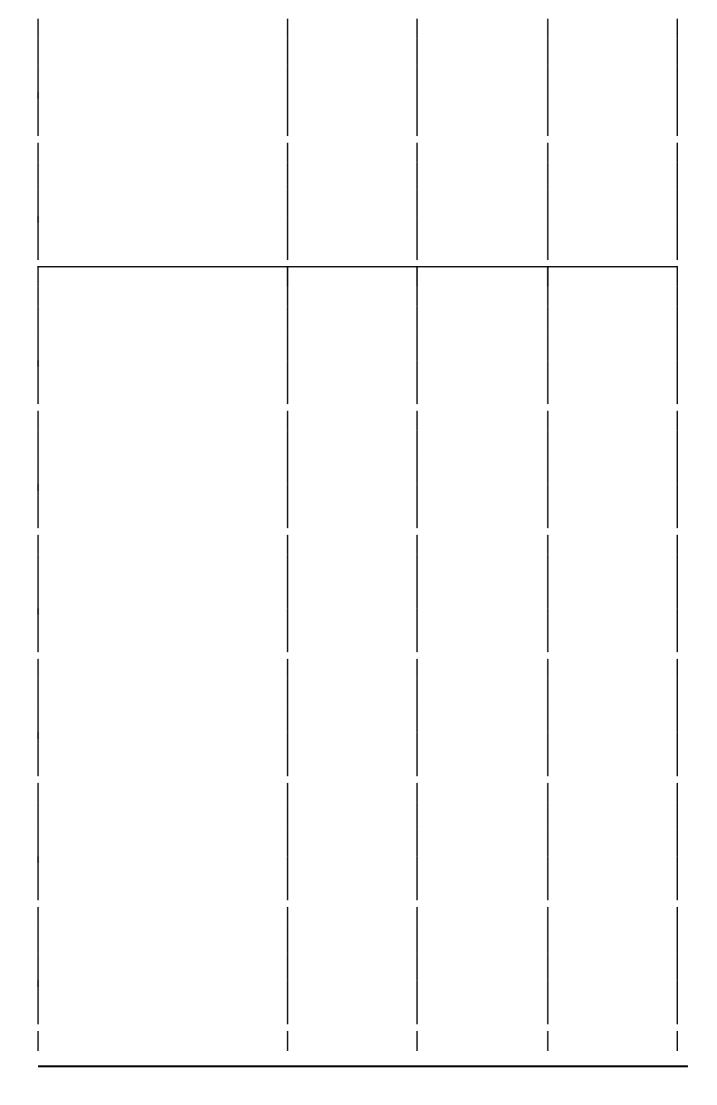


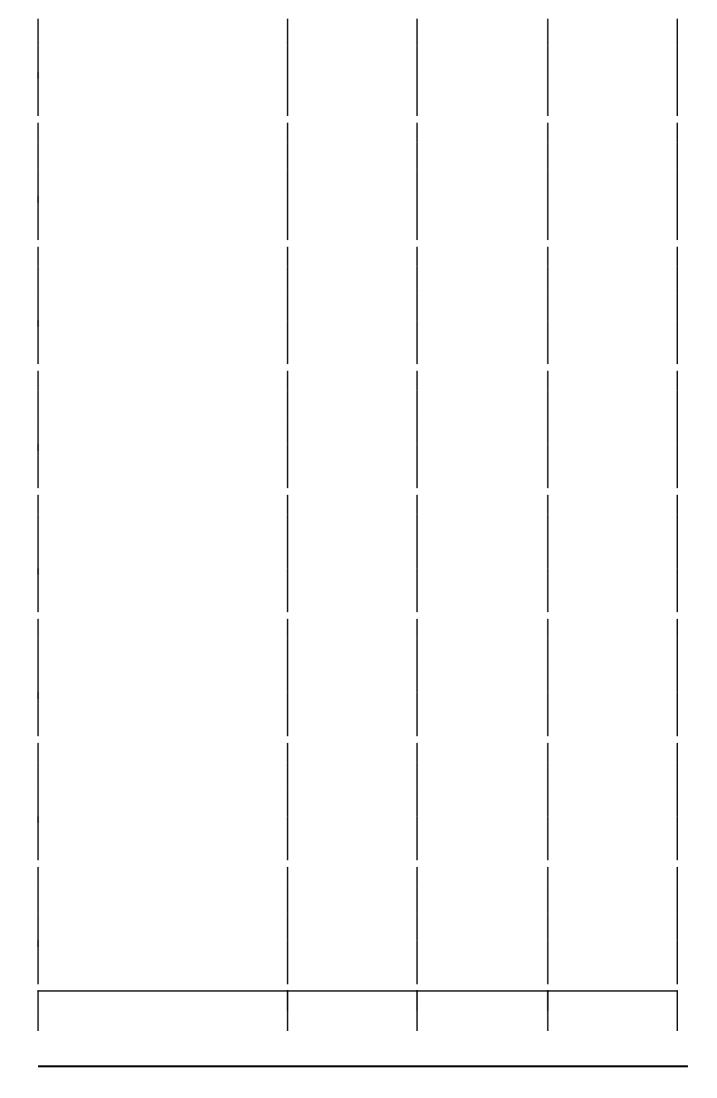


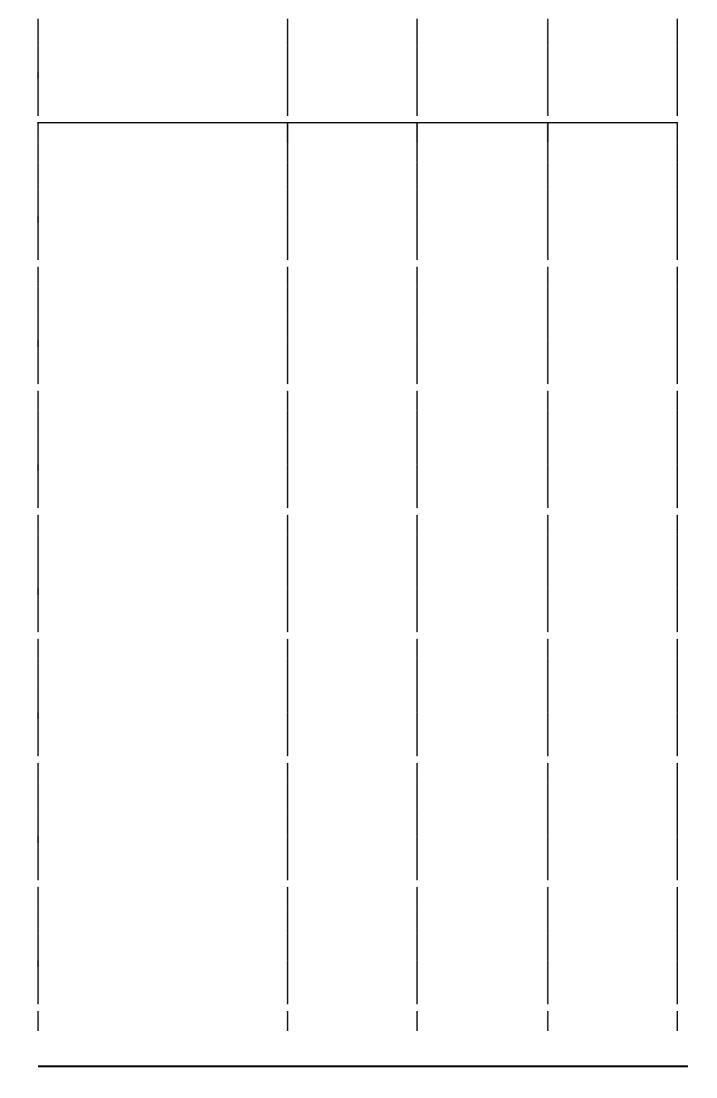


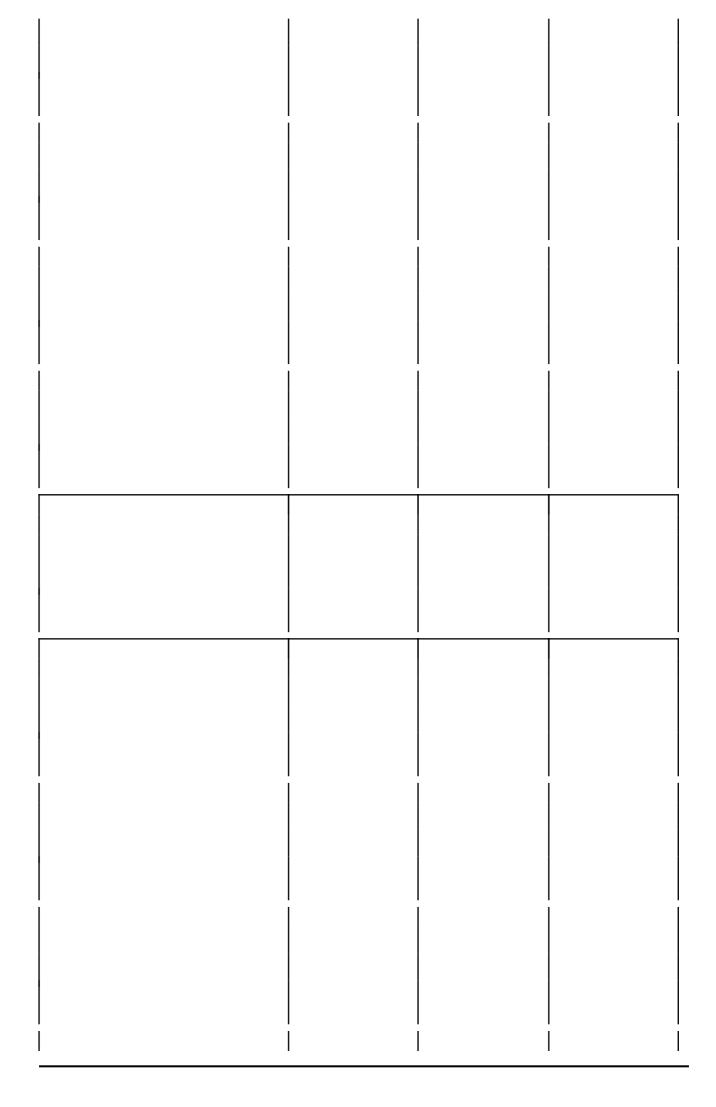


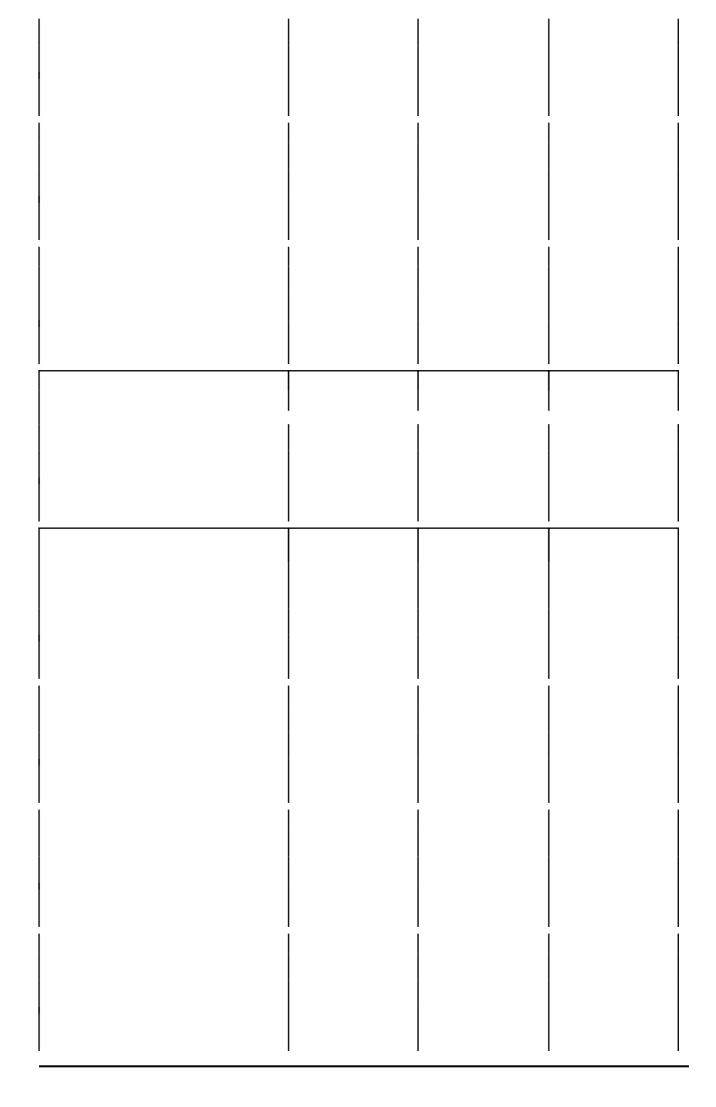


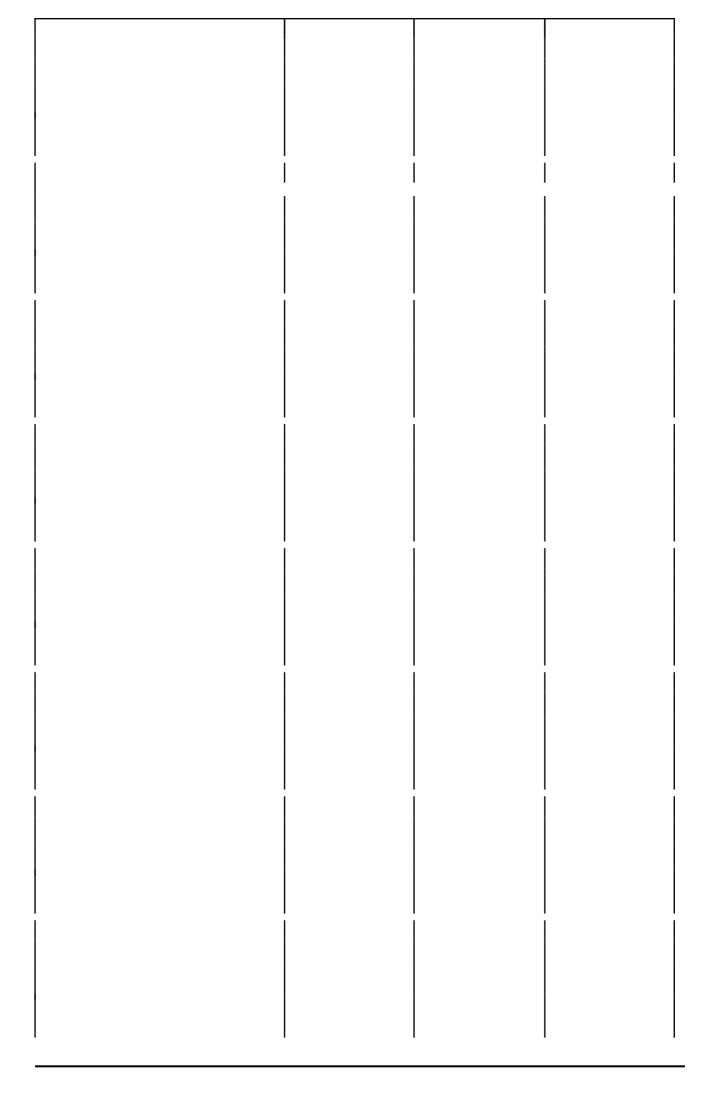


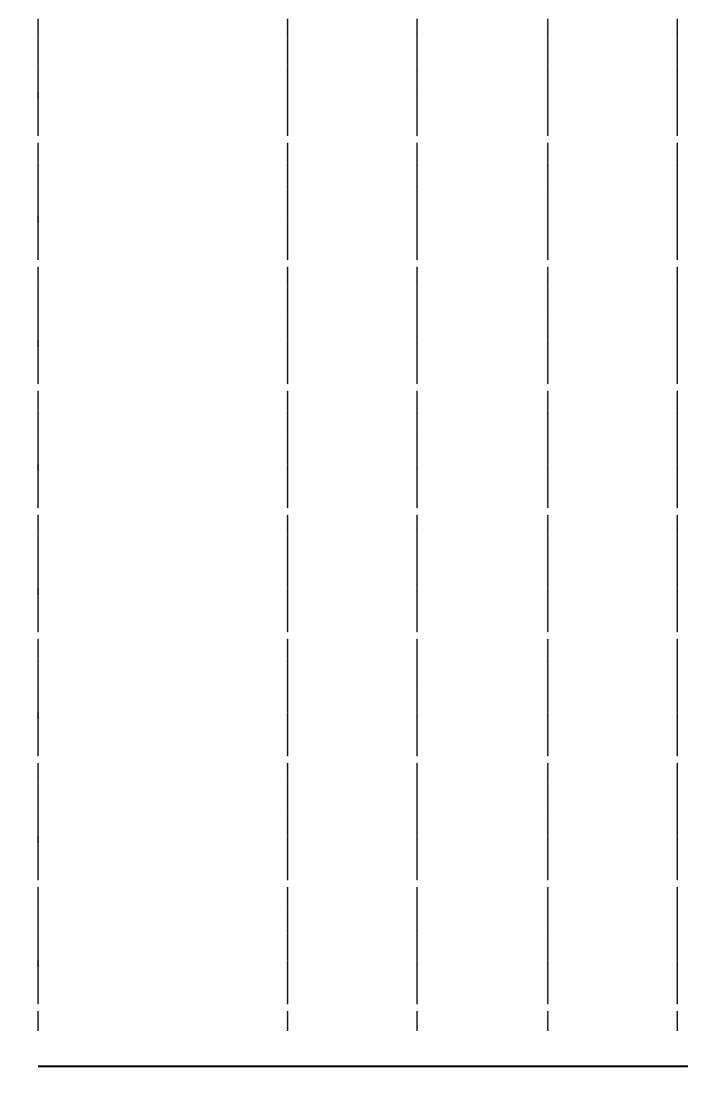


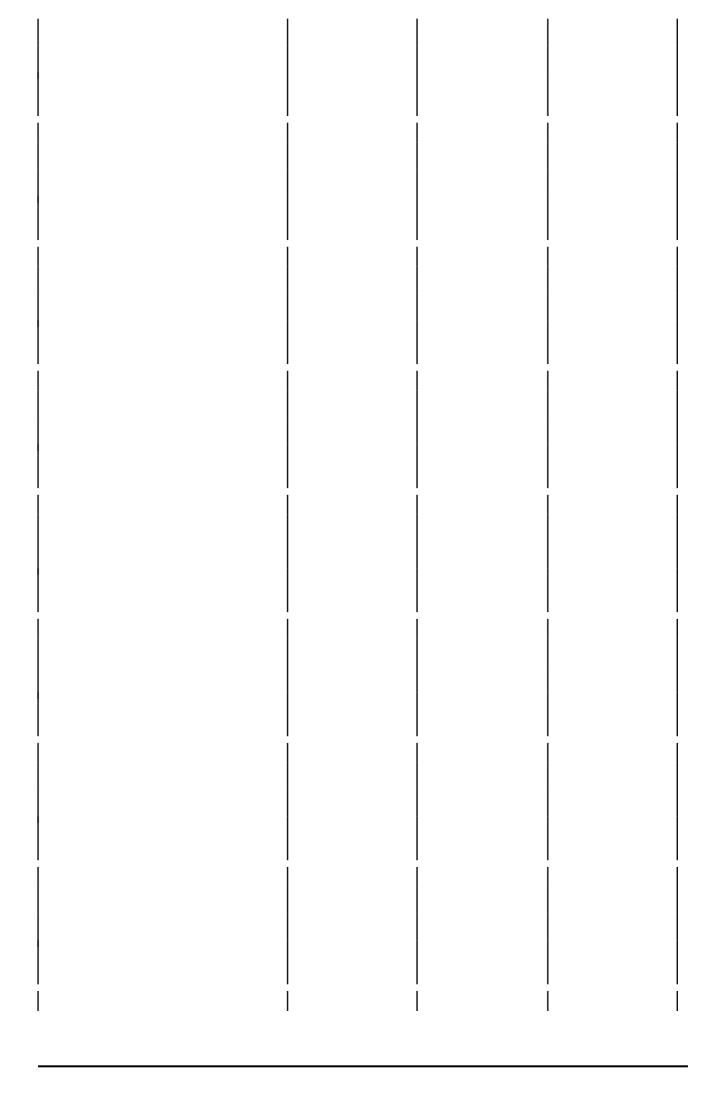




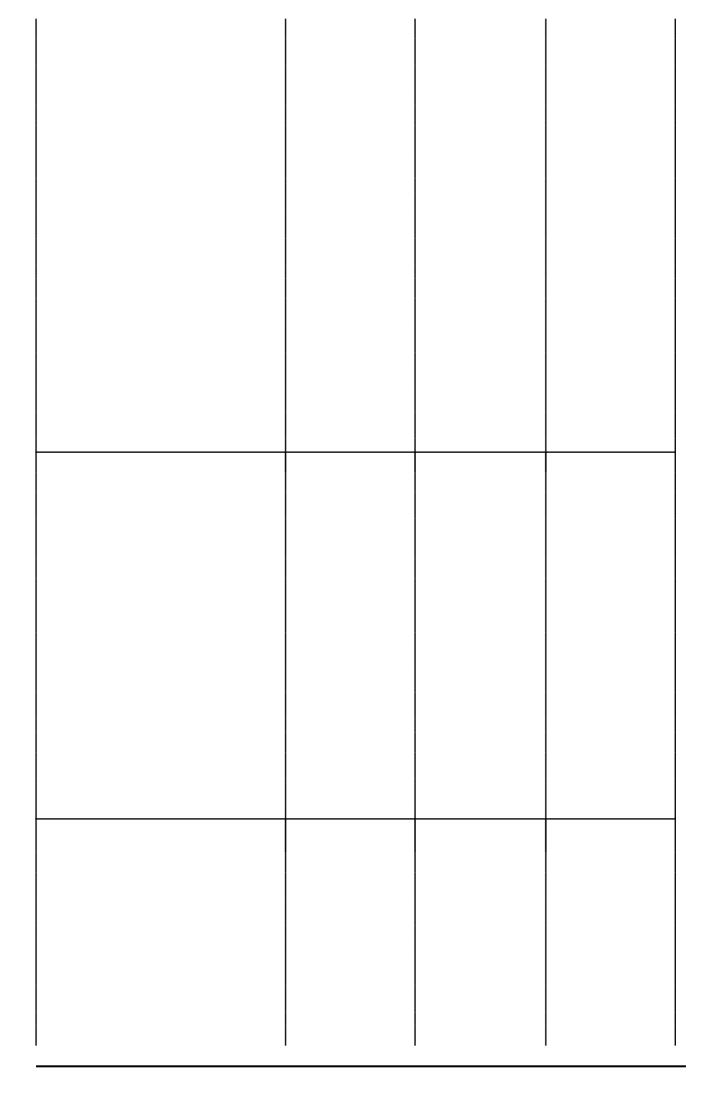


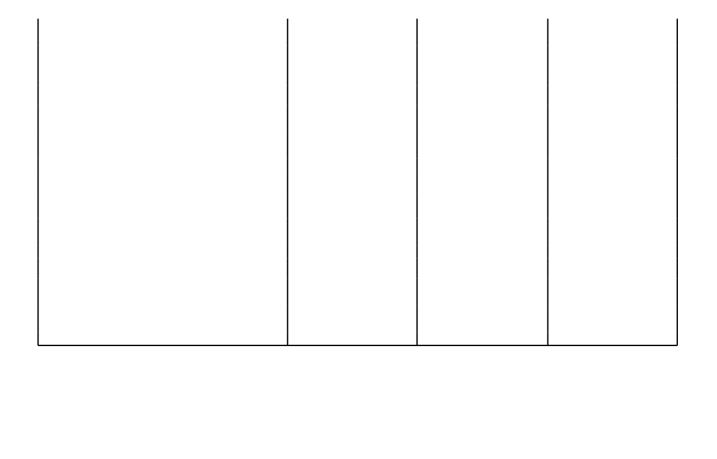






| Non-serious adverse events | | |
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More information

Substantial protocol amendments (globally)

| Date | Amendment |
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Interruptions (globally)

Limitations and caveats