

Disclosure Results

Cyramza

European Marketing Authorisation Holder: Eli Lilly Nederland B.V.

| phase | total complete by 31 July 2016 | un-evaluable | evaluable | disclosed in 12 month timeframe | disclosed percentage at 12 months | complete by 31 July 2016 | disclosed at 31 July 2016 | disclosure percentage at 31st July 2016 |
|--------------|--------------------------------------|--------------|-----------|---------------------------------------|---|-----------------------------|------------------------------|--|
| phase I & II | 26 | 1 | 25 | 24 | 96% | 25 | 24 | 96% |
| phase III | 6 | 0 | 6 | 6 | 100% | 6 | 6 | 100% |
| phase IV | 0 | 0 | 0 | 0 | | 0 | 0 | |
| other | 0 | 0 | 0 | 0 | | 0 | 0 | |
| total | 32 | 1 | 31 | 30 | 97% | 31 | 30 | 97% |

Footnote (company communication): One phase II trial (conducted in the US and Canada) remained undisclosed at 31 July 2016 but has since been published.

Key to columns in table above:

total complete by 31 July 2016 = total number of company-sponsored trials identified which were completed by 31 July 2016

unevaluable = trials with completion date within the last 12 months, or key dates missing - excluded from the analysis

evaluable = trials with all criteria present including dates; hence the base number of trials evaluated for the assessment

disclosed in 12 month timeframe = evaluable trials which were disclosed within the target 12 months*

disclosed percentage at 12 months = proportion of evaluable trials which were disclosed within 12 months*

complete before 31 July 2016 = number of evaluable trials completed before 31 July 2016

disclosed at 31 July 2016 = number of evaluable trials with results disclosed by 31 July 2016

disclosure percentage at 31 July 2016 = proportion of evaluable trials which were disclosed by 31 July 2016

[* 12 months measured from the later of: the first date of regulatory approval (in Europe or the US) or the trial completion date]