# Key FSP Medical Writing Experience

| Sponsor | Document Types | Regions Supported | Duration of Relationship | Employees |
|---|---|---|---|---|
| Sponsor 1 | CSRs, narratives, basic results/FDAAA, QC/QR-only | U.S., EU, CN | 5 yrs | Up to 60 |
| Sponsor 2 | Safety aggregate reports, risk management profiles | U.S., EU | 2+ yrs | Up to 17 |
| Sponsor 3 | Submission documents, narratives, protocols | U.S. | 2 yrs | 10+ |
| Sponsor 4 | CSRs, narratives, submission documents | AU, IN | 2 yrs | 34 |
| Sponsor 5 | Multiple regulatory documents (CSRs, BDs, RPs, etc), submissions, publications, slide sets | U.S., EU, Japan | 8 yrs | Up to 50 |
| Sponsor 6 | Multiple regulatory document types and submission modules, safety aggregate reports, publications, patient narratives, CM&C documents, standalone editing and QC, SOP technical writing, CTR support, lay language docs | U.S., EU, Japan, China, Australia | 12 yrs | Up to 117 |
| Sponsor 7 | CSRs, narratives, MedInfo, basic results/FDAAA, standalone QC | U.S., EU | 6 yrs | Up to 41 |
| Sponsor 8 | Safety aggregate reports | U.S., EU | 2 yrs | 6 |
| Sponsor 9 | Clinical trial lay summaries | EU | 2 yrs | Up to 4 |

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