

JESSICA LEWIS

(470) 559-6313

jessicaelewis22@gmail.com

CAREER SUMMARY

An adept Medical Writer with medical information expertise supporting the oncology therapeutic area. Translated complex scientific data to create clinical dossiers and Q&A documents that supported successful launches for various pharmaceutical and biotechnology accounts across a wide range of therapeutic areas, including oncology and rare diseases. Contributed to pharmacovigilance activities by accurately documenting and reporting safety information to support regulatory compliance. Crafted medically accurate promotional materials and deliverables in market access and medical communications for large global pharmaceutical and medical device companies, including Sanofi, Dexcom, Johnson & Johnson, Baxter, and Olympus.

INDUSTRY RELEVANT EXPERIENCES

Precision AQ| Atlanta, GA

Scientific Associate/ Medical Writer

March 2025- Present

- Translate complex scientific data into payer value propositions, clinical dossiers, health system formulary access kits, and Q&A documents to support successful launches across a wide range of therapeutic areas, including oncology, rare diseases, and diabetes.
- Collaborate with Sanofi's cross-functional teams during the medical, legal, and regulatory (MLR) review process to ensure promotional and educational materials are scientifically accurate, legally compliant, and meet FDA regulatory standards.
- Conduct 200+ comprehensive literature searches and ongoing surveillance of scientific publications to identify relevant clinical data and potential adverse event reports.
- Research and analyze cancer clinical trials and real-world evidence to provide a competitive strategic landscape and develop value stories for brand positioning and strategy for cross-functional teams and external clients.
- Collaborate with Sanofi's Medical Affairs, Regulatory, and Pharmacovigilance teams to ensure scientific accuracy and alignment.
- Spearhead multiple launches and develop high-quality scientific and medical information materials, meeting 100% of established deadlines to support the launch and timely drug approval of a multimodal anti-CD38 therapy delivered via an on-body delivery system for multiple myeloma.
- Maintain 100% compliance with internal standard operating procedures (SOPs) and regulatory standards for medical information responses.

Experts Ink| Atlanta, GA
Medical Writer/ Medical Editor

November 2023- January 2025

- Conducted rigorous literature reviews to ensure the accuracy and validity of medical information, enhancing credibility.
- Translated complex medical jargon into clear, easy-to-understand content for a diverse audience, including healthcare professionals, patients, and corporate stakeholders.
- Critically analyzed and summarized 100+ scientific publications to support medical information responses.

Edanz| Atlanta, GA
Freelance Medical Writer

November 2022- September 2023

- Spearheaded 100+ projects to streamline the company's medical information database, resulting in increased audience engagement.
- Optimized medical content for adaptability across various target audiences, including healthcare professionals, patients, and internal stakeholders.
- Supported Scientific Communications initiatives by synthesizing data from clinical studies into concise, written deliverables.

HCL America| Kissimmee, FL
Regulatory Medical Writer

February 2021- June 2022

- Attended meetings with Baxter's clinical product team to provide evaluation input for updating Clinical Evaluation Reports (CERs) and Post-Marketing Surveillance (PMS) documents.
- Collaborated with Johnson & Johnson's Regulatory team and developed multiple clinical documents, including Clinical Study Reports (CSRs), clinical protocols, and investigator brochures, to support FDA and EMA submissions.
- Recorded data in the safety database with 100% accuracy and completed all applicable tracking of incoming safety information related to Baxter's medical devices.
- Crafted compelling narratives for Baxter's global brand strategy, enhancing consistency across international markets and increasing brand recognition and engagement by 80%.
- Demonstrated leadership by mentoring associate medical writers and contributing to team knowledge-sharing initiatives, thereby enhancing overall team productivity and document quality.
- Applied strong scientific and analytical expertise to interpret complex clinical, scientific, and technical data, synthesizing information from multiple sources into clear, compliant regulatory documents for the oncology therapeutic area.

PROFESSIONAL EXPERIENCE

Medicine Shoppe Pharmacy| Kissimmee, FL Registered Pharmacist

June 2020- December 2020

- Responded to unsolicited medical inquiries from healthcare professionals and patients with evidence-based information regarding various drug products available in the pharmacy.
- Performed Academy of Managed Care Pharmacy (AMCP) practices by implementing Medication Therapy Management (MTM) to assess medication costs, monitor adherence, and document adverse events.
- Collaborated with pharmacy technicians to prepare and dispense an average of 150–300 prescriptions daily while communicating with healthcare providers and patients to ensure the safe use of medications.

RESEARCH EXPERIENCES

University of Florida, College of Medicine| Gainesville, FL Graduate Research Assistant

April 2017- May 2020

Research topic: USF1 in the regulation of breast cancer progression

- Provided scientific support at national and international medical conferences, including the American Association for Cancer Research (AACR) Annual Meeting, engaging with healthcare professionals and scientists.
- Collaborated with a cross-functional team to develop a mouse model of human breast cancer to investigate changes in tumor weight and volume resulting from overexpression of a gene in a cell line.

EDUCATION

ISI Language School Tokyo-Takadanobaba Campus

Japanese Language (N3-N2)

July 2022- June 2023

University of Florida, College of Medicine

MS Thesis in Molecular Cell Biology/ Medical Sciences

April 2017- May 2020

University of Pittsburgh School of Pharmacy

Doctor of Pharmacy, PharmD

August 2010- December 2014

SKILLS

- **Medical Writing:** Promotional materials, Market Access writing, Medical Information writing skills, Clinical study reports, Abstracts, Value dossiers, MSL slide decks, Literature evaluation, Investigator brochures, Executive summaries, Clinical protocols
- **Soft Skills:** Client-oriented, Attention-to-detail, Critical thinking, Ability to meet tight deadlines, Time management, Cross-functional team working skills, Business agility, Problem-solving, Resource management, Mentoring, Great communication skills
- **Technical Skills/ Software:** Microsoft Office, PubMed, EndNote, Grammarly, ChatGPT, Veeva, GraphPad Prism
- **Languages:** English (native), Japanese (business level), Korean (basic level)
- **Medical knowledge & Compliance:** European Union Medical Device Regulation (EU-MDR), FDA regulations, AMA style guide, Medical terminology, Multiple therapeutic areas (especially Oncology and Neurology), Biostatistics

ADDITIONAL INFORMATION

- **Licensure:** Pharmacist License (RPh), Florida (PS 54638) —issued 2015-present
- **Certifications:** American Medical Writers Association (AMWA) Essential Skills Certificate (2021), Japanese Language Proficiency Test, Level N3 (2020), Area of Concentration in Alzheimer's Research (2014), Continuing Education Recognition in Chemotherapy (2014), Pharmacy-Based Immunization Delivery (2012)
- **Professional Affiliations:** Active Member of AMWA, Cheeky Scientist Medical Writing Association

REFERENCES

Available upon request