

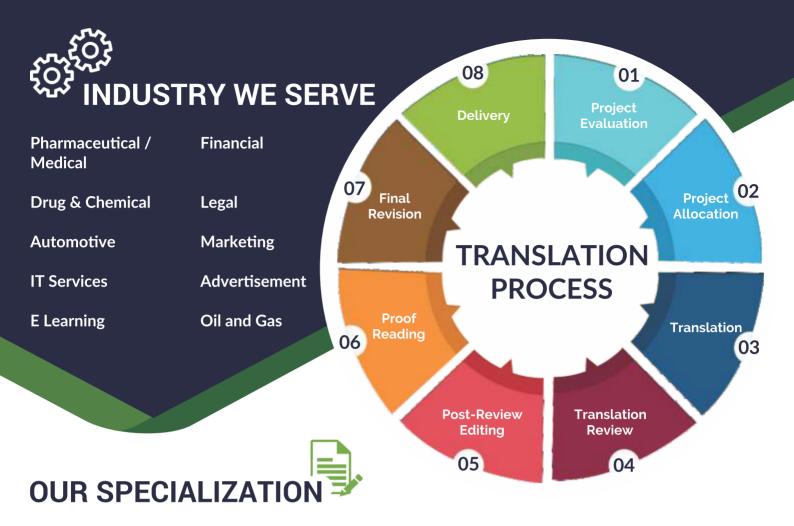
ISO 9001: 2015 CERTIFIED CO.



Neoplus Translation is one of the foremost names in the field of translation. Translation being a vast field, consisting of varied forms, our expertise lies in the **Pharmaceutical translation** as the founder Nishit Shah is **Qualified Pharmacist (M. Pharm in Pharmaceutics)**

Neoplus Translation, established in 2012, is an establishment with a specific criterion wherein we provide the professional translation services in the various sectors to the clients as per their requirements. We have an entire team of qualified linguists who can cater to the client's needs just as they would need it.

Our company believes in providing best services with an entire plethora of languages covering Europe (EU), CIS, Asian and Emerging market. Accuracy, relevancy and articulation are the key elements. We mandate client satisfaction which is why we have been chosen for various government initiatives and our clientele base is a testament to our success.



Our Specialization is Pharmaceutical/Medical Translation as the owner himself is Pharmacist (M. Pharm in Pharmaceutics) with team of Pharmacist and Regulatory Affairs Expert.

Pharmaceutical Translation Service



As an international pharmaceutical translation provider, Neoplus guides Pharmaceutical companies through the regulatory requirements that demonstrate the opportunities for pharmaceutical sales across the European Union and in its member states. Our job is helping you — Marketing Authorization Holders for new drug products — translate your pharmaceutical product (and win business) in more than 20 official languages of the European Union.

NEOPLUS Translation brings unparalleled value by

- Working with native-speaker translators certified in clinical translation for each Member State;
- Independent medical consultants and regulatory experts who provide in-country reviews for linguistic and technical accuracy following latest QRD.



Translate In the recent years there has been a phenomenal growth in the pharmaceutical industry of Commonwealth of Independent States (CIS). All the CIS nations have been focusing on building trade relations with various countries especially in the field of pharmaceuticals and biotechnology.

Language is one major barrier to conduct business in CIS countries as they accept Pharmaceutical or Regulatory documents only in Russian and their native language. NEOPLUS Translation has in-house native certified translators working in our Indian office which has a thorough Pharmaceutical background.

We have translated 150+ Drug Dossiers and DMFs till now for top notch Pharmaceutical companies in order for submission in CIS countries.



EMERGING MARKET

Emerging market extends from Latin America to Asia and Africa. NEOPLUS Translation helps Pharmaceutical companies to build an agile organization by providing translation service in native language of the intended country which facilitates the need and provides them a relative ease in doing business.

We cover all Languages for Emerging market and the major ones: Spanish, Portuguese, Arabic, African languages, Vietnamese, Chinese, Indonesian, Korean, Turkish, etc.

We have translated about 150+ Drug Dossiers/ DMF in Spanish language for submission in Mexico and Latin American countries, 100+ in French language for submission in Francophone countries, 50+ in Chinese language for submission in China



MEDICAL DEVICE DOCUMENT TRANSLATION

Selling your medical device globally requires adaptation as per foreign markets. NEOPLUS Translation specializes in the translation and localization of medical and surgical device documents, and helps to ensure that the translated documents follow international and national standards technically, linguistically and culturally.



Medical Documents We Translate

- Drug Dossiers (Module 1 to Module 5)
- Drug Master File (DMF) (3.2.S)
- Patient Information Leaflet (PIL)
- Summary of Product Characteristics (SPC)
- Package Insert (PI)
- Normative Documents
- Material Safety Data Sheet (MSDS)
- Site Master File (SMF)
- PSUR (Periodic Safety Update Report)

- Clinical Protocols
- Clinical Trials
- Contracts
- Pharmacological Studies
- Product Labels
- Protocols
- Informed Consent Forms
- Investigator Brochures
- SAE and SOP Procedures

- Corporate Websites and Portals
- Instruction Manuals
- Device Patents
- Operator's Manuals
- Regulatory Guidelines
- Pharmacovigilance documents
- Regulatory Audit Reports
- Master Batch Records
- Scientific Journal Articles

Our Prestigious Clients





















































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