

MEDICAL & PHARMACEUTICAL TRANSLATION

Specialist translation and review for life-sciences content

Spanish (Spain & Latin America) · Brazilian Portuguese · European languages · into English

15

years focused exclusively on medical and pharmaceutical content

2

linguists on every project: a specialist translator and a second reviewing linguist

17100

ISO-aligned workflow: our process is aligned with ISO 17100

For pharmaceutical and biotech sponsors, CROs, clinical research sites, regulatory consultancies and translation agency partners.

WHO WE ARE

A specialist partner for medical translation

Latin Link was founded by scientific translators. For 15 years we have worked exclusively on medical and pharmaceutical content, helping life-sciences organisations communicate accurately with regulators, investigators and patients.

Our core service is **translation and review**. Every project is translated by a native specialist medical linguist and reviewed by a second linguist, then quality-checked before delivery. A dedicated team (account manager, project managers and medical linguists) looks after every client.

15 years
exclusive focus on medical and pharmaceutical content

Translation + review
every project, by two qualified linguists

5+ years
minimum field experience for our medical linguists, who also have clinical experience

Same-day
jobs available, with capacity to scale for Brazilian Portuguese

EXPERIENCE YOU CAN CHECK

Pfizer
We support Pfizer with translation into English.

The BMJ
We cover the BMJ's *Good Medical Practice* content every month.

CAPABILITIES SUMMARY

Core service	Translation by a native specialist medical linguist + review by a second linguist + final QA
Languages	Spanish (Spain), Spanish (Latin America), Brazilian Portuguese, European languages, into English
Content	Clinical, regulatory & CMC, safety, patient-facing, medical device, publications and medical communications
Regulatory	Documentation for ANVISA, COFEPRIS, INVIMA, DIGEMID and ANMAT submissions
Quality	Process aligned with ISO 17100; workflows follow industry best practice
Tools	Trados and memoQ; client portal for sending files and paying for projects
Clients	Sponsors, CROs, research sites, regulatory consultancies, agency partners

TRANSLATION & REVIEW

Two linguists on every project, as standard

Our core service is built around one principle: no translation reaches you without being checked by a second specialist. It's the safeguard regulators, ethics committees and patients rely on.

01

Brief & file preparation

Files arrive via our client portal. Your project manager confirms scope and deadline and prepares glossaries and reference material.

02

Specialist translation

A native medical linguist with clinical experience translates into their mother tongue, within their field.

03

Independent review

A second qualified linguist reviews the translation against the source for accuracy, terminology, completeness and style.

04

Final QA

Checks on numbers, units, terminology consistency, formatting and completeness before release.

05

Delivery

Delivered on time in your required format, ready for your submission, site or publication.

ISO 17100 ALIGNED

Aligned with ISO 17100

ISO 17100 is the international standard for translation services. Our workflow is aligned with it, and our processes follow industry best practice.

WHAT THE STANDARD EXPECTS	HOW WE WORK
Qualified, competent translators	Native specialist medical linguists with clinical experience and a minimum of 5 years' experience in the field
Revision by a second, different linguist	Every project is reviewed by a second qualified linguist
Managed projects and client communication	Dedicated project managers and an account manager for every client
Verification before delivery	Final QA on completeness, consistency and formatting

WHAT WE TRANSLATE

Documents across the product lifecycle

From study start-up to post-marketing safety, we translate and review the documents your clinical, regulatory and medical teams depend on.

Development → Submission → Labelling → Safety → Communications

Clinical development

Clinical operations · CROs · sites

- Clinical study protocols, synopses and amendments
- Investigator's Brochures (IBs)
- Informed consent forms and patient information sheets
- Clinical study reports (CSRs)
- Case report forms (CRFs)
- Ethics committee and site correspondence

Regulatory & CMC

Regulatory affairs · labelling

- Regulatory dossier modules, including CMC/quality
- Summaries of product characteristics (SmPCs)
- Patient information leaflets (PILs)
- Labelling and package inserts
- Material safety data sheets (MSDS)
- Documents for Latin American submissions

Safety & pharmacovigilance

Drug safety · PV teams

- Pharmacovigilance reports
- Periodic safety reports
- Adverse event reports and case narratives
- Hospital and medical reports

Patient-facing

Clinical operations · patient engagement

- Patient questionnaires and clinical outcome assessments (COAs)
- Patient information and education materials
- Patient-facing trial materials

Medical devices

Regulatory · product teams

- Instructions for use (IFUs) and directions for use (DFUs)
- Device labelling and packaging text
- User manuals and safety information

Publications & medical communications

Medical affairs · publishers

- Journal articles, manuscripts and abstracts
- *Good Medical Practice* content for the BMJ, monthly
- Medical education and brochures
- Healthcare websites and marketing materials



04

Languages & regulatory coverage

OUR LANGUAGES

ES — **Spanish — Spain**
European Spanish for EU studies, submissions and patient materials.

LA — **Spanish — Latin America**
Regional Spanish for Mexico, Colombia, Peru, Argentina and beyond.

PT — **Brazilian Portuguese**
Capacity to scale for high-volume programmes and same-day jobs.

EU — **European languages**
Specialist medical linguists across European language pairs.

EN — **Into English**
UK and US English for source documents, publications and regulators.

LATIN AMERICAN REGULATORY SUBMISSIONS

We translate documentation for sponsors and regulatory consultancies preparing submissions to the region's health authorities.

ANVISA	**Brazil**	Agência Nacional de Vigilância Sanitária
COFEPRIS	**Mexico**	Comisión Federal para la Protección contra Riesgos Sanitarios
INVIMA	**Colombia**	Instituto Nacional de Vigilancia de Medicamentos y Alimentos
DIGEMID	**Peru**	Dirección General de Medicamentos, Insumos y Drogas
ANMAT	**Argentina**	Administración Nacional de Medicamentos, Alimentos y Tecnología Médica

One team for multi-country studies. Protocols, ICFs, labelling and dossiers for Latin American programmes handled by one team, with consistent terminology across countries.

OUR TEAM

A dedicated team behind every project

Project managers and specialist medical linguists working together, with an account manager for every client, so you always know who is looking after your work.

Account manager

A dedicated point of contact for every client, who knows your projects, preferences and terminology.

Project managers

Experienced, multilingual project managers who plan, coordinate and deliver complex projects on time.

Medical linguists

Native specialist translators and reviewers with clinical experience, working only within their field.

OUR MEDICAL LINGUISTS HAVE

- Clinical experience
- A minimum of 5 years' experience in the field
- Native-language translation only, within their specialism
- Degree-level education
- An utmost respect for deadlines



WHO WE WORK WITH

Pharma & biotech sponsors	Clinical, regulatory and safety content across the lifecycle.
CROs	Responsive capacity for multi-country study documentation.
Clinical research sites	Patient-facing and site documents, translated and reviewed.
Regulatory consultancies	Submission documentation for Latin American authorities.
Translation agency partners	Specialist medical Spanish, Brazilian Portuguese and into-English capacity.

WORKING WITH US

Simple to start, reliable to scale

Client portal

Send files and pay for projects online through our client portal, convenient for direct clients and agency partners alike.

Technology

Experts in Trados and memoQ, with translation memories and glossaries for consistent terminology. We adapt to your preferred CAT tool, and we have mainstream desktop publishing software for print-ready files.

Turnaround & scale

Same-day jobs available, and capacity to scale for large Brazilian Portuguese programmes, with a no-nonsense attitude to deadlines.

Confidentiality

Client documents are handled confidentially and shared only with the team working on your project. We are happy to work under your NDA.

FOR A FAST QUOTE, SEND US

1. Your source files
2. Target language(s) and market
3. Your deadline
4. Glossaries or reference material

WHY LATIN LINK

Six reasons teams choose us

01
Specialised
15 years exclusively in medical and pharmaceutical content.

02
Reviewed, always
Every project translated and then reviewed by a second linguist.

03
Clinically experienced
Linguists with clinical experience and 5+ years in the field.

04
Latin America experts
Spanish for Spain and Latin America, Brazilian Portuguese and regional regulators.

05
Dedicated team
An account manager and project managers who know your work.

06
Easy to work with
Client portal, flexible tools and same-day capacity.

LATIN LINK

START A PROJECT

Let's talk about your next project

Whether you're a sponsor, CRO, research site, regulatory consultancy or agency partner, send us your files for a free, no-obligation quote.

01 · Share your files

Upload through our client portal or send them by email.

02 · Receive your quote

We confirm scope, languages and delivery date.

03 · Translated & reviewed

Delivered on time, reviewed by a second linguist.

Luke Sewell

Client Service Director

EMAIL
luke.sewell@latinlink.com

TELEPHONE
+44 (0)20 3286 4122

MOBILE
+44 (0)7795 083485

WEB
www.latinlink.com

ADDRESS
12 Hammersmith Grove
London W6 7AP, UK

CLIENT PORTAL
Send files and pay for projects online