Trench Rossi Watanabe.

LIFE SCIENCES

At the forefront of the Life Sciences industry for more than 60 years, we deeply understand the challenges faced by clients

We have a **multidisciplinary team highly specialized in several legal practices**, including regulatory, intellectual property, public law, compliance, civil, criminal, transactional and tax. We advise **pharmaceutical**, **medical device**, **and other companies in the industry** to enter the market, grow and achieve excellent results in a **wide range of matters**, while always anticipating trends.

We have extensive experience in **handling complex corporate transactions**, **regulatory procedures**, **drug pricing approval**, **drafting technology licensing and other agreements of the sector** and defining **market access strategies**. We help clients deal with conflicting **regulatory regimes** and the related impacts of expansion, manufacturing and distribution strategies. We help companies file and protect **patents**, conduct clinical trials to gain approval for new drugs and products, and comply with rules relating to the collection, use and treatment of **patient data**.

We have an excellent standing with the National Health Surveillance Agency (ANVISA); the National Research Ethics Commission (CONEP); the Drug Market Regulation Chamber (CMED); the National Supplementary Health Agency (ANS); the Ministry of Agriculture, Livestock and Food Supply (MAPA); the State Health Secretariats; and the municipal health authorities.

We also have a **strong presence in litigation** involving product liability and unfair competition. We offer support in **compliance issues and digital health**. We also participate in the **drafting of bills** and regulations affecting the industry, and we help our clients **navigate the complex tax regime** applicable to the healthcare sector to optimize their results.

How we can help

- Advising on issues related to regulation of the health sector
- Advising on Good Manufacturing Practices Certification
- Consulting in Productive Development Partnerships (PDP), Technological Purchase Orders (ETECs), and other models of innovative partnerships in healthcare
- Technology transfer agreements
- Drafting and reviewing agreements related to manufacturing, distribution, co-marketing, among others
- Advising on issues involving clinical trials before Research Ethics Committees and the National Research Ethics Commission (CEPs and CONEP)
- Public procurement advisory and relationship with the Ministry of Health
- Administrative defenses and appeals against ANVISA, MAPA and local health authorities
- Advising on market access (issues) involving CMED, CONITEC and ANS
- Advising on corporate transactions involving the sector
- Healthcare compliance program consulting
- Advising on issues involving the certification of regulated products before the National Institute of Metrology, Quality and Technology (INMETRO)
- Representation in litigation involving patents and trademarks of medicines, health products and other regulated products

Awards and recognitions

IP STARS from Managing IP

LEADERS LEAGUE

Chambers AND PARTNERS

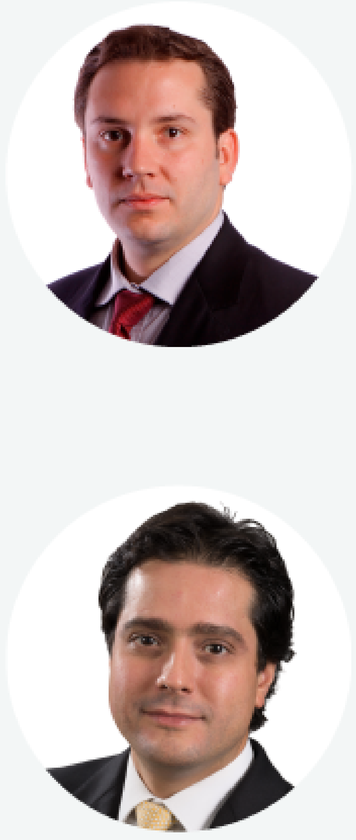
The LEGAL 500

análise ADVOCACIA 500

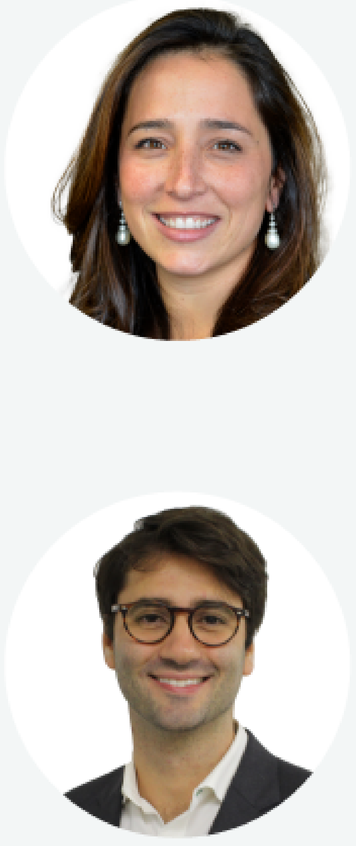
Core team

Henrique Frizzo
Partner
henrique.frizzo@trenchrossi.com

Marcela Trigo
Partner
marcela.trigo@trenchrossi.com



Mauricio Pacheco
Partner
mauricio.pacheco@trenchrossi.com



Felipe Zaltman
Partner
felipe.zaltman@trenchrossi.com

Carla Moraes
Associate
carla.moraes @trenchrossi.com

for more information, use the QRCode or click here.

Trench Rossi Watanabe.

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