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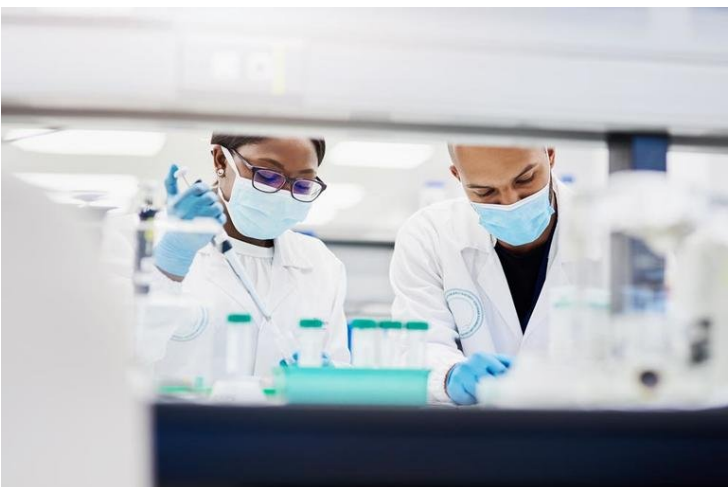
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HRV Pharma and Shodhana Labs forge CDMO alliance to fast-track next-gen API development

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India's first virtual API platform enters longevity biology & precision medicine



HRV Global Life Sciences (HRV Pharma), India's first integrated Virtual API and Pharmtech company, has announced a comprehensive, multi-year Contract Development & Manufacturing partnership with Shodhana Laboratories, a Hyderabad-based USFDA-aligned API manufacturer, to co-develop, manufacture and supply a curated pipeline of high-science APIs, including pharmaceutical-grade API's primarily used in longevity biology, backed by a US Drug Master File (DMF).

This makes HRV the holder of the first and only active Indian DMF in this product as a pharmaceutical API. The collaboration brings together HRV's global market access across 50+ countries, AI-enabled Virtual API platform and regulatory DMF engine, with Shodhana's deep process chemistry, high-purity synthesis expertise and GMP infrastructure creating a seamless development-to-commercialisation pathway for molecules at the frontier of precision medicine.

The partnership has already launched with five active development programs, with both organizations jointly managing regulatory strategy and documentation under a structured, milestone-gated framework.

Under the multi-year agreement, both companies will jointly execute:

- **Development and US DMF filing for pharmaceutical-grade API** — a master coenzyme governing cellular energy, DNA repair and aging, now emerging as a regulated therapeutic API in oncology supportive care, neurodegeneration and longevity medicine.
- **A pipeline of five high-science APIs** spanning CNS, rare/orphan disease, and metabolic/neurological API's — with partner anchor end-customers across the USA and EU
- **End-to-end GMP manufacturing, CMC documentation, stability studies and validation** aligned to USFDA, EMA, PMDA and ANVISA standards
- **Global regulatory DMF filings across US, EU, LATAM, MENA & APAC** with HRV managing all regulatory interactions and expenses
- **A unified quality, compliance and audit-readiness framework** for the highest tier of regulated market access