



Tobacco Harm Reduction Solutions

DVCR offers end-to-end clinical and regulatory support from study design through CSR submission. We operate a 100-bed clinical unit that is backed by a team with deep industry experience with both tobacco and nicotine products.

PMTA-Relevant Clinical Evidence

- ✓ Abuse Liability
- ✓ PK/PD
- ✓ Biomarkers of Exposure
- ✓ Biomarkers of Potential Harm
- ✓ Product use Topography
- ✓ Bioequivalence
- ✓ Oral Health Endpoints
- ✓ Disease Related Endpoints
- ✓ Longitudinal Safety Studies



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Full-Service CRO

- ✓ 100-bed Phase I Unit
- ✓ Multi-Site Trials
- ✓ Project Management
- ✓ Data Management
- ✓ Biostatistics & Programming
- ✓ Medical Writing
- ✓ Clinical Monitoring
- ✓ Medical Monitoring
- ✓ Bioanalysis
- ✓ PK/PD Analysis & Modeling

We offer PMTA and FDA regulatory support focused on clinical study design, execution and submission-ready data packages.

Smoking Rooms

Housed within our facility are seven indoor exposure rooms, including two large and five individual rooms.

- ✓ Enables PK/AL data collection while volunteers use study products
- ✓ Dedicated ad libitum and controlled-use environment
- ✓ Allows direct observation of product use behavior
- ✓ Design allows different group sizes
- ✓ Accommodates different product types