Elenco NON esaustivo degli acronimi utilizzati nell'industria farmaceutica

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Acronimo	Definizione
AIC	Autorizzazione Immissione in Commercio
AIFA	Agenzia Italiana del Farmaco
API	Active Pharmaceutical Ingredient
CAPA	Corrective and Preventive Action
ASMF	Active Substance Master Files
CMC	Chemistry, Manufacturing and Controls
CP	Centralized Procedure
CRC	Clinical Research Coordinator
CRO	Contract Research Organisation
CSR	Clinical Study Report
CTD	Common Technical Dossier
CTC	Clinical Trial Certificate
DMF	Drug Master File
EDC	Electronic Data Capture
FDA	Food and Drug Administration
GCP	Good Clinical Practice
GDP	Good Distribution Practice
GLP	Good Laboratory Practice
GMP	Good Manufacturing Practice
GPvP	Good Pharmacovigilance Practise
GVP	Good Vigilance Practises
HMPC	Committee on Herbal Medicinal Products
ICDRA	International Conference of Drug Regulatory Authorities
IDE	Investigational Device Exemption
ISO	International Organization for Standardization
LoA	Letter of Access
MD	Medical Device
NB	Notified Body

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Acronimo	Definizione
NBF	Norme di Buona Fabbricazione
NDA	New Drug Application
NDS	New Drug Submission
NIH	National Institutes of Health
NPA	National Pharmaceutical Association
OD	Orphan Drug
PIL	Product Information Label
PIM	Product Information Management
PRAC	Pharmacovigilance Risk Assessement Committee
PSMF	Pharmacovigilance Sistem Master File
PSUR	Product Safety Update Report
PVAR	Preliminary Variation Assesment Report
RMS	Reference Member State
SCADA	Supervisory Control and Data Acquisition
SPC	Supplementary Protection Certificate
QA	Quality Assurance
QC	Quality Control
QOS	Quality Overall Summary
QP	Qualified Person
QPPV	Qualified Person Pharmacovigilance
QRD	Quality Review of Documents
REACH	Registration, Evaluation, Authorisation and Restriction of Chemicals
R&D	Research and Development
RMP	Risk Management Plan
SMF	Site Master File
SOP	Standard Operating Procedure