



API's  
INTERMEDIATES  
CUSTOM SYNTHESIS  
CONTRACT R&D  
cGMP FACILITIES



**CORAL**  
DRUGS PRIVATE LIMITED



## an adventure called science

Edwin Hubble once wrote “Equipped with his five senses, man explores the universe around him and calls the adventure Science”. A love of this adventure is what drives us at Coral Drugs.

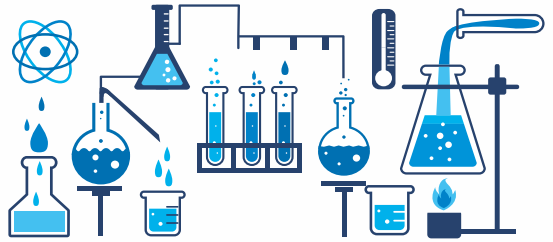
Everything we do at Coral is derived from our love of science, which demands rigour, discipline and dedication. In return, we get satisfaction in watching our products make lives better for people all over the world, everyday.

The knowledge that the quality of our products is trusted globally is a huge responsibility on Coral as a company. It pushes us to be ever vigilant and meticulous in all that we do, so that our future may live up to our past.

At Coral we believe that we are not just making APIs, we are creating a world where people can live safe and healthy lives.



**C**oral Drugs was incorporated in 1993, with a vision to make the world a better place through our love of science and began production in 1996. We started with processes for APIs and intermediates for anti-cancer drugs and steroids. Today, what started as just another pharmaceutical manufacturing unit is a supplier of APIs to major pharmaceutical companies around the world.



The journey from those early days till now has been long and fruitful. On the way, we've spent countless nights obsessing over particle sizes, types of machinery and delivering schedules and ironed out many details. We have seen our products become an indispensable part of many major drugs and companies worldwide. Thanks to the quality of our APIs, we are effecting change on a global level and helping people all over the world live better lives everyday.

coral /  
a passion



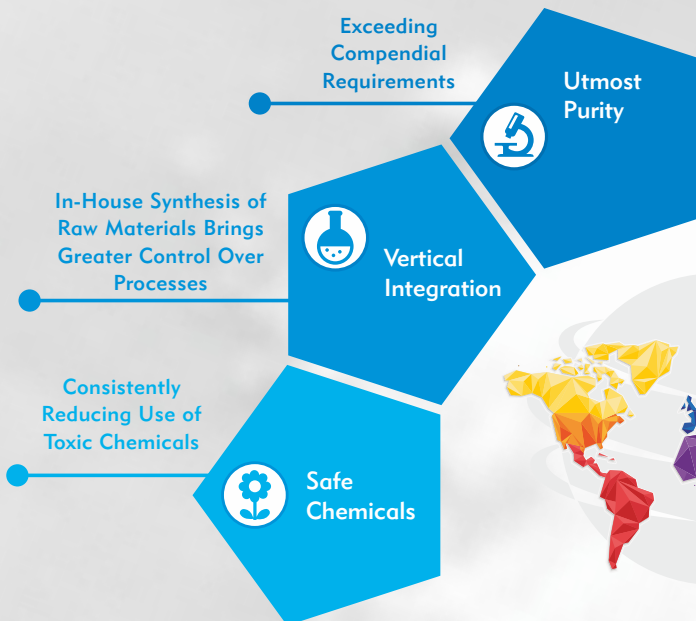
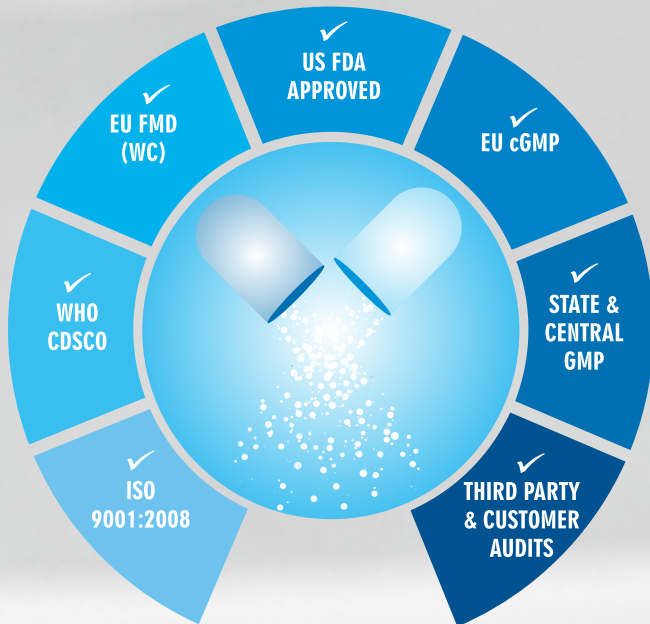
**quality is a surety** ♥

Science demands purity of the highest order, and quality is paramount. This is why, we at Coral, take immense pride in the quality of our products and go to great lengths to ensure that exacting standards are met 100% of the time.

Each of our process have a validated SOP which we strictly follow. Audits, both internal and external, are regularly conducted to evaluate and maintain the health of our system. Our units - which comply with cGMP and GLP and are WHO and ISO certified - have been audited by the United States Food and Drug Administration (US FDA), other regulatory bodies and by our partners themselves.

Our team is fully trained at induction. This is augmented by on-the-job training, which is followed by regular checks and validations to ensure adherence to policies and procedures. Science does not tolerate inferior quality, and neither do we.

**“ We take immense pride in the quality of our products and go to great lengths to ensure our exacting standards are met 100% of the time ”**



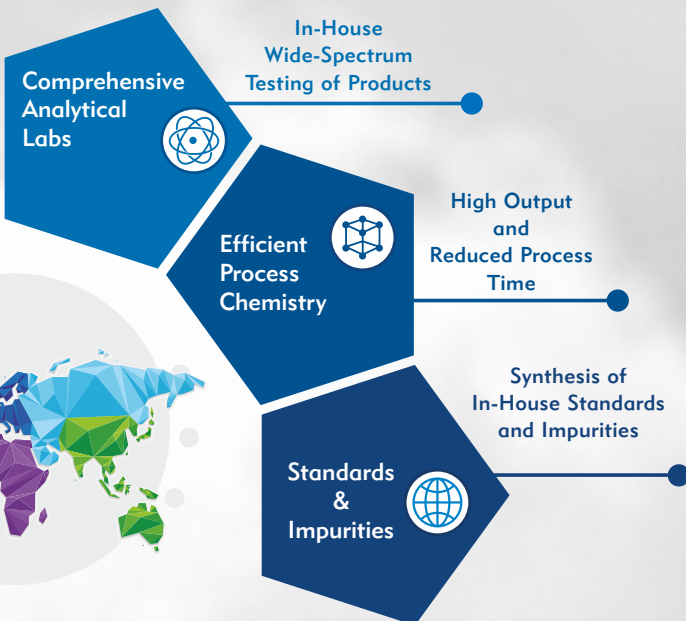
Our manufacturing facilities are well equipped for practically any challenge

SIZE DOES MATTER ♥

The efficacy of every API that we produce is important and is dependent to a large extension particle size. We realise that particle size can impact the bioavailability and the dosage form of every API, as well as dissolution, absorption rate and content uniformity. At Coral, through a two-tiered process, we ensure to deliver the correct particle size every time.

Firstly, we ensure that the correct particle sizes are produced. Whether it is controlled by our chemical processes or a variety of milling and grinding techniques, our manufacturing facilities are well equipped for practically any challenge. Once the product is milled and ready, the next step is to measure the particles to ensure that they are of the correct size.

We believe that the most important aspect of any particle size analysis is the development of a rugged and reproducible method of analysis. Our team spends countless hours developing methods that minimize particle agglomeration by using a range of compatible surfactant and non-soluble dispersant media.



size does matter



## how we create value

At Coral, our core internal attributes are aligned in a manner that provides a customer experience that is unparalleled in our industry.

Internally: Our value proposition starts with a deep **Focus** towards the therapeutic areas that we operate in. We combine this with the **Experience** of our various quality teams to deliver a truly **Efficient Synthesis** of our products.

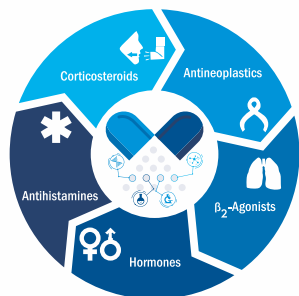
Further: Our **Dedication** to cGMP manufacturing results in very high **Production Standards** and through our **Continuous Research**, we frequently opt for **Environmentally** safer synthesis.

Finally: Our **Global Approach** helps us deliver quality products to our client's world over.

Such operations and structure help us deliver quality products **Consistently**. The results of which are evident to our clients who have come to expect a high level of **Quality** and **Customization** to fit their specific needs.

Moreover, our **Timely Deliveries** of materials and support, help our clients meet their own goals with ease. We test ourselves both, internally and externally, by continuing to participate in Global audits (**Quality Approvals**) and Regulatory filing around the world.





## Our Active Pharmaceutical Ingredients (APIs)\*

### CORTICOSTEROIDS

- BUDESONIDE US DMF CEP DMF
- CICLESONIDE ▲
- CLOBETASOL PROPIONATE
- FLUNISOLIDE
- DEFLAZACORT ▲
- FLUOCINOLONE ACETONIDE
- FLUOCINONIDE
- FLUTICASONE FUROATE US DMF DMF
- FLUTICASONE PROPIONATE US DMF CEP DMF
- ISOFLUPREDONE ACETATE ▲
- MOMETASONE FUROATE US DMF DMF
- MOMETASONE FUROATE MONOHYDRATE US DMF DMF
- TRIAMCINOLONE DMF
- TRIAMCINOLONE ACETONIDE US DMF CEP DMF
- TRIAMCINOLONE HEXACETONIDE DMF

### ANTIHISTAMINES

- AZELASTINE HYDROCHLORIDE DMF

### LABA (Long Acting Beta Agonists)

- FORMOTEROL FUMARATE DIHYDRATE US DMF CEP DMF
- INDACATEROL MALEATE
- SALMETEROL XINAFOATE ▲

### ANTINEOPLASTICS

- DANAZOL DMF
- ESTRAMUSTINE SODIUM PHOSPHATE DMF
- EXEMESTANE DMF
- FLUTAMIDE CEP DMF

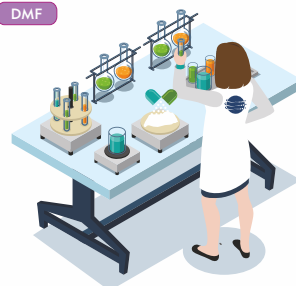
### HORMONES

- MEDROXYPROGESTERONE ACETATE
- NORETHISTERONE DMF
- NORETHISTERONE ACETATE DMF

▲ Tech Pack Under Development

\*Products currently covered by valid US patents are offered for R&D use in accordance with 35 USC 271(e)+A13(1). It is the buyer's responsibility to verify the patent position in markets of intended use before contracting with us. Any patent infringement and resulting liability is solely at the buyer's risk.

Ask us about your Custom Synthesis or cGMP Contract Manufacturing needs





**CORAL**  
DRUGS PRIVATE LIMITED

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we  
♥  
science

