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Descriptions_tools = [
{
    "name": "annotate_open_reading_frames",
    "description": (
        "Find all Open Reading Frames (ORFs) in a DNA sequence using Biopython. "
        "Scans the forward strand and (optionally) the reverse-complement strand, "
        "returning ORF coordinates that meet a minimum nucleotide length."
    ),
    "required_parameters": [
        {
            "name": "sequence",
            "type": "str",
            "default": None,
            "description": (
                "DNA sequence to analyze. Should be provided as a raw string of nucleotides "
                "(typically A/C/G/T; clarify whether ambiguous bases like N are allowed)."
            ),
        },
        {
            "name": "min_length",
            "type": "int",
            "default": None,
            "description": (
                "Minimum ORF length in nucleotides (not amino acids). ORFs shorter than this are
                excluded."
            )
        }
    ]
}
]

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    ),
  },
],
"optional_parameters": [
  {
    "name": "search_reverse",
    "type": "bool",
    "default": False,
    "description": (
      "If True, also scan the reverse-complement strand. This typically increases runtime "
      "by ~2x on the same input length."
    ),
  },
  {
    "name": "filter_subsets",
    "type": "bool",
    "default": False,
    "description": (
      "If True, filter out ORFs that share the same end position but have a later start "
      "(i.e., remove subset ORFs nested at the same stop site)."
    ),
  },
],
"hardware_requirements": {
  "device": "cpu_only",
  "notes": "No GPU required. Performance depends mainly on sequence length and CPU single-
thread speed.",

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    },
    "time_complexity": {
        "assumptions": (
            "Report wall-clock latency for a typical input size (e.g., 5–10 kb plasmid, 1 Mb contig, etc.),
"
            "and specify CPU model + threads. Include cold-start vs warm-start if Biopython import
dominates."
        ),
        "latency_seconds": {"n1": None, "n2": None, "n10": None}, # doc-writer fills with
benchmarks
    },
},
{
    "name": "annotate_plasmid",
    "description": (
        "Annotate a DNA sequence by invoking pLannotate via its command-line interface. "
        "Typically used for plasmid annotation (features, CDS, common parts) and supports "
        "circular sequences."
    ),
    "required_parameters": [
        {
            "name": "sequence",
            "type": "str",
            "default": None,
            "description": (
                "DNA sequence to annotate (raw nucleotide string). Clarify expected length range "
                "and whether FASTA headers/newlines are accepted."
            )
        }
    ]
}

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    ),
  }
],
"optional_parameters": [
  {
    "name": "is_circular",
    "type": "bool",
    "default": True,
    "description": (
      "If True, treat the sequence as circular during annotation (important for plasmids). "
      "If False, treat as linear DNA."
    ),
  },
],
"hardware_requirements": {
  "device": "cpu_only",
  "notes": (
    "No GPU required. Requires pLannotate installed and accessible in PATH. "
    "May need substantial RAM depending on database/index usage."
  ),
},
"time_complexity": {
  "assumptions": (
    "Benchmark includes CLI startup + any database/index loading. Provide times for typical
    plasmid sizes "
    "(e.g., 3–20 kb) and specify CPU + storage type (network FS vs local SSD can matter)."
  ),

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"latency_seconds": {"n1": None, "n2": None, "n10": None}, # doc-writer fills with
benchmarks
},
}
]
```

Name

What it is:

The unique function/tool identifier.

Complete it by:

- Matching the exact function or CLI wrapper name in code.
- Keeping it stable across versions (avoid renaming).
- Using snake_case and being explicit (e.g., `annotate_open_reading_frames`, not `orf_tool`).

Why it matters:

Agents and pipelines call this string directly — any mismatch breaks execution.

description

What it is:

A high-level explanation of what the tool does, suggestions on when to use the tool and explanation about how good the tool is compared to the others.

Complete it by:

- Stating input → processing → output clearly.
- Naming the underlying library/algorithm (Biopython, CLI, model, etc.).
- Mentioning key behavior assumptions (strand search, circular DNA, etc.).
- Keeping it concise (2–4 sentences max).

Why it matters:

Helps humans and agents choose the correct tool without reading code.

required_parameters

What it is:

Inputs that must always be provided by the agent for the tool.

Complete it by (for each parameter):

- name → exact argument name in code
- type → strict type (str, int, bool, etc.)
- description → what it means + units + format
- default → usually None, but if a value is said by the GitHub to be fixed set the default value

Also specify:

- expected format (FASTA vs raw string)
- units (nt vs aa)
- allowed ranges
- constraints or validation rules

Why it matters:

Prevents silent failures and incorrect inputs.

optional_parameters**What it is:**

Inputs that modify behavior but aren't required.

Complete it by (for each parameter):

- real default value (must match code)
- clear explanation of what changes when enabled/disabled
- note performance or memory impact if applicable

Why it matters:

Prevents ambiguous behavior and hidden runtime costs.

Hardware_requirements**What it is:**

The compute environment needed to run the tool.

Complete it by specifying:

- cpu_only / gpu_optional / gpu_required

Example:

- CPU only

Why it matters:

Lets schedulers, clusters, or agents route jobs correctly.

Time_complexity

What it is:

Real-world runtime measurements.

Complete it by:

- Measuring wall-clock time (not Big-O math)
- Stating assumptions:
 - hardware used
 - input size
 - threads/batching
- reporting:
 - time for 1 run
 - time for 2 runs
 - time for 10 runs

Example:

- $1 \rightarrow 0.4s$
- $2 \rightarrow 0.7s$
- $10 \rightarrow 3.2s$

Why it matters:

Agents can plan batching, parallelization, and scheduling accurately.

Outputs

What it is:

What the tool returns.

Complete it by specifying:

- data type (list, dict, dataframe, file path, JSON)
- schema/fields
- units/format
- example output

Why it matters:

Downstream tools must parse this reliably.

Failure_modes**What it is:**

Common errors and how to handle them.

Complete it by listing:

- error condition
- likely cause
- recommended fix

Example:

- empty sequence → invalid input → validate before call

Why it matters:

Prevents silent crashes in pipelines.

We also will require requirements for each tool and function call:

**Dependencies should be in a separate .txt file
(requirements_Function_name.txt)****What it is:**

External libraries, binaries, or models required.

Complete it by specifying:

- package names + versions
- installation method

- whether system binary or Python package

Why it matters:

Ensures reproducibility across machines/HPC nodes.