

MAP chemotherapy is a standard treatment regimen for osteosarcoma and consists of three drugs: **methotrexate**, **doxorubicin** (also called Adriamycin), and **cisplatin**[157](#).

- **Purpose:** Used primarily for high-grade osteosarcoma (a type of bone cancer), MAP is given to shrink tumors before surgery and reduce recurrence risks after surgery, often with curative intent[14](#).
- **How it is given:**
 - Administration is typically in **cycles**, with each cycle followed by a rest period to allow for recovery[18](#).
 - Patients often receive **2 cycles before surgery** and **4 cycles after surgery**, with each cycle lasting about **35 days**[18](#).
 - Drugs are delivered intravenously through a central line, PICC line, or portacath[1](#).
- **Efficacy:** MAP remains the **preferred regimen for osteosarcoma** based on current evidence. Meta-analyses have not shown significant survival differences between MAP and more intensive combinations (such as adding ifosfamide and etoposide), but MAP is associated with **lower rates of serious toxicities**[56](#).
- **Toxicity and side effects:**
 - The most common serious side effects are **neutropenia** (low white blood cells), **thrombocytopenia** (low platelets), **anemia**, and **febrile neutropenia** (fever with low white cells)[256](#).
 - Compared to more intensive regimens, MAP causes these toxicities less frequently[56](#).
- **Key drug-specific toxicities:**
 - *Methotrexate*: kidney toxicity, mucositis.
 - *Doxorubicin*: heart toxicity (risk increases with cumulative doses)[3](#).
 - *Cisplatin*: kidney and hearing damage, nerve toxicity.
- **Standard of care:** There is strong expert consensus from major trials (such as EURAMOS-1) that MAP is the standard treatment for resectable osteosarcoma, though no other chemotherapy regimens have shown clear superiority[35](#).

Important caveats:

- While MAP is the global standard for osteosarcoma, individual patient factors, tumor response, and local protocols may lead to regimen adjustments.
- The regimen is intensive and requires close monitoring for toxicity.

MAP, therefore, is a cornerstone of osteosarcoma therapy due to its balance of efficacy and manageable toxicity compared to alternatives⁵⁶.

1. <https://www.cancerresearchuk.org/about-cancer/treatment/drugs/map>
2. <https://pmc.ncbi.nlm.nih.gov/articles/PMC5052459/>
3. <https://www.eviq.org.au/medical-oncology/sarcoma/bone-sarcoma/1901-osteosarcoma-map-methotrexate-doxorubicin>
4. <https://www.chemoexperts.com/map-methotrexate-doxorubicin-cisplatin-sarcoma.html>
5. <https://pmc.ncbi.nlm.nih.gov/articles/PMC6531127/>
6. [https://journals.lww.com/md-journal/fulltext/2019/05100/methotrexate, doxorubicin, and cisplatinum regimen.90.aspx](https://journals.lww.com/md-journal/fulltext/2019/05100/methotrexate,_doxorubicin,_and_cisplatinum_regimen.90.aspx)
7. <https://www.cancer.org/cancer/types/osteosarcoma/treating/chemotherapy.html>
8. <https://www.eviq.org.au/medical-oncology/sarcoma/bone-sarcoma/1901-osteosarcoma-map-methotrexate-doxorubicin/patient-information>
9. <https://ascopubs.org/doi/10.1200/JCO.2014.60.0734>