

<b>Characteristic</b>	<b>Examples of relevant evidence</b>
<b>1. Is Electrophilic or Can Be Metabolically Activated</b>	Parent compound or metabolite with an electrophilic structure (e.g., epoxide, quinone, etc), formation of DNA and protein adducts.
<b>2. Is Genotoxic</b>	DNA damage (DNA strand breaks, DNA-protein cross-links, unscheduled DNA synthesis), intercalation, gene mutations, cytogenetic changes (e.g., chromosome aberrations, micronuclei).
<b>3. Alters DNA repair or causes genomic instability</b>	Alterations of DNA replication or repair (e.g., topoisomerase II, base-excision or double-strand break repair)
<b>4. Induces Epigenetic Alterations</b>	DNA methylation, histone modification, microRNA expression
<b>5. Induces Oxidative Stress</b>	Oxygen radicals, oxidative stress, oxidative damage to macromolecules (e.g., DNA, lipids)
<b>6. Induces chronic inflammation</b>	Elevated white blood cells, myeloperoxidase activity, altered cytokine and/or chemokine production
<b>7. Is Immunosuppressive</b>	Decreased immunosurveillance, immune system dysfunction
<b>8. Modulates receptor-mediated effects</b>	Receptor in/activation (e.g., ER, PPAR, AhR) or modulation of exogenous ligands (including hormones)
<b>9. Causes Immortalization</b>	Inhibition of senescence, cell transformation
<b>10. Alters cell proliferation, cell death or nutrient supply</b>	Increased proliferation, decreased apoptosis, changes in growth factors, energetics and signaling pathways related to cellular replication or cell cycle control, angiogenesis

FIGURE 1