zvou je nejen výběr vhodných léků, ale i management nových nežádoucích účinků. Další, neméně palčivou problematikou, je i chybějící úhrada u některých léků. Navzdory prvotním slibným výsledkům cílené terapie se zdá být nepravděpodobné, že zasažení jediné patologické signální dráhy by mohlo vést k dlouhodobému vyléčení. Odhalení molekulárních mechanismů leukemogeneze a rezistence může přispět k lepšímu pochopení signálních drah a vývoji účinnější terapie. Další intenzivní výzkum a vývoj nových léků nadále probíhá, aplikace jejich výsledků do klinické praxe je tak příslibem pro zlepšení prognózy a dlouhodobých léčebných výsledků AML.

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