

Commentary

is impro isation in the techniq\(\subseteq es \) of cell biolog

Comp⊠ter scientists and statisticians ha e also contrib⊠ted for the impro ed image processing and anal sis. e s nergistic e orts of the cell biologists and the other professions ha e enabled the is ali ation

of cells ar tracking path a s in the lie cells at greater resolstion. e interactions bet een the di erent organelles co de be seen b capt⊠ring three dimensional color footage of light sheet Øorescence microscop combined ith confocal microscop. is has opened

Microscop has re ollowioni ed the a lord thlio o -Baran CdtaCs d ildne and the a lord thlio o -Baran CdtaCs d ildne and the lord by the lo and test the biological h pothesis. Se eral of the cellar processes incloding mitosis and meiosis ere decoded being these ad anced microscopic is ali ations. Understanding of inter molec ar interactions has great rele ance in the pharmace tical sector. On an o erall basis the imaging capabilit at s\square cell\square are le els has open ne a en\subseteq es and far more attracti e for the researchers ho are interested in cell biolog and its applications.

> Genetic screening and editing ha e enabled the scientists the reengineer the cells and cellar processes and daring these process the in ol ement of mathematical, statistical and comparer science ha e been phenomenal. Die to ad ancements in the technolog sach as genetic engineering for e perimental cell biolog large scale e periments ere possible and this has resetted in generation of hage olames of comple and d namic data that needed to be anal ed and interpreted. e di ersit and the magnit⊠de of the molec@ar and ph siological processes necessitated the anal sis of the data generated to deri ed logical interpretations. Genetic reprogramming is enabling the scientists to reengineer the cells state. that the detect an abnormalit in the earl stage. e techniq e is also enabling regenerati e medicine and cell regeneration and tiss De reconstr⊠ctions. e compatation pro ess, machine learning and modeling has enabled anal sis of ariables associated ith tho ands of genes and gene combination e ects. All the technologies ha e facilitated s nthesis and prod\(\mathbb{\omega}\)ction of biological therape\(\mathbb{\omega}\)tics s\(\mathbb{\omega}\)ch as monoclonal antibodies for ario\(\mathbb{O} \) therape\(\mathbb{D} \) tic applications. Cell biolog is increasingl becoming more rele ant to health science that

Technological Frontiers in Cell Biology

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en mes \(\mathbb{D}\) nctioned. E er since the microscopic de ices ere in ented there has been constant impro ement in the technolog and in fact the de elopment of high resolation Dorescence microscope has resalted in conferring Nobel Pri e in Chemistr in the ear 2014. In the ear 2017 the pri e ent to the de elopment of cr o-electron microscope. e Anderl ing orking principles of these microscopes di er a great deal. In orescence microscop the orescence molec's are sed to light the target proteins,, cells, and other cell ar components, and e en

allo s the biologists to is ali e the li ing samples on a real time basis. Ho e er, there as a limitation that isible light cannot di erentiate components closer than 200 nanometers. erefore electron microscope as de eloped in order to achie e greater resol\(\mathbb{D}\)tion. B\(\mathbb{D}\)t ho e er this techniq De req Dired the ac Dom that limited its Doe for is Dali ing li e erefore of techniq es of cr stallograph ere replaced ith the indi id⊠al or combined ⊠sage of these microscopes.

Wide eld microscope f⊠nctions on the intense light so⊠rce to ill minate the specimen. Confocal microscope as de eloped that Sees a pin hole to ill minate the point of interest. is enables remo al of backgroand orescence and oat of focas elds and hence a higher resolation and contrast as obtained in e ect. Light sheet microscop forctions b scanning the samples based on a er thin plane of laser light and this techniq e allo s real-time tracking of the li ing cells and tiss Des. Similarl the str Det Dred ill Demination microscop also allo s the magni ed is ali ation of the li e cells. e most ad anced electron cr o-microscope has the potential to re eal that atomic str⊠ct⊠res of biomolec⊠res s⊠ch a large protein. is has man ad antages o er X-ra cr stallograph beca\(\mathbb{Z}\)ee ith this techniq\(\mathbb{Z}\)es the samples req\(\mathbb{Z}\)ired to be ed and a cr stal form.

e er before.